


**Gamma-1 and Gamma-3 Anti-Human CD23 Monoclonal
Antibodies and Use Thereof as Therapeutics**

Cross Reference to Related Application

5  This application is a continuation-in-part of U.S.
Serial No. 08/803,085, filed February 20, 1997.

Field of the Invention

10 The present invention relates to monoclonal
antibodies which specifically bind human CD23, the low
affinity receptor for IgE (FceRII/CD23), and contain
either a human gamma-1 or human gamma-3 constant domain,
and their usage as therapeutic agents.

Background of the Invention

15 IgE is a member of the immunoglobulin family that
mediates allergic responses such as asthma, food aller-
gies, type 1 hypersensitivity and the familiar sinus
inflammation allergic rhinitis and conjunctivitis, and
as a result, causes widespread suffering throughout the
general population. IgE is secreted by, and expressed
on the surface of, B-cells. IgE synthesized by B-cells
20 can be anchored in the B-cell membrane by a short
transmembrane domain linked to the mature IgE sequence.
Membrane and secreted versions of IgE are formed in the
same cell by differential splicing of the IgE RNA
transcript.

25 IgE also can be bound to B-cells (and T cells,
monocytes, Langerhans cells, follicular dendritic cells,
natural killer cells, eosinophils and platelets) through
its Fc region to a low affinity IgE receptor (FceRII,
hereafter "FCEL", and to mast cells and basophils
30 through its Fc region to a high affinity IgE receptor
FceRI, (hereinafter "FCEH"). The low affinity IgE

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receptor is generally referred to in the literature as CD23.

Upon exposure of a mammal to an allergen, antigen presenting cells process the antigen for presentation to helper T cells. These helper T cells secrete cytokines such as IL-4 which assist B-cells to undergo clonal amplification and secrete more allergen-specific IgE. This newly synthesized IgE in turn is released into the circulation where it binds to mast cells and basophils through the high affinity receptor on their cell surface. Such mast cells and basophils are thereby sensitized to the specific allergen. The next exposure to the same allergen causes binding to specific IgE on the surface of mast cells, and basophils, thereby cross-linking the FcεRI on these cells and thus activating their release of histamine and other factors which are responsible for clinical hypersensitivity and anaphylaxis.

The art has reported antibodies capable of binding to FcεL (CD23)-bound IgE but not IgE bound to FcεH (see, for example, WO 89/00138 and U.S. Patent 4,940,782). These antibodies are disclosed to be clinically advantageous because they bind to IgE which is bound to the low affinity receptor (FcεL) or to circulating IgE's, but do not bind to IgE bound to the high affinity receptor (FcεH). Therefore, these antibodies will not activate mast cells or basophils.

Moreover, anti-CD23 antibodies have been reported to have potential as therapeutics, e.g., for the treatment of allergic disorders, inflammatory diseases, and autoimmune diseases. For example, Bonnefoy et al., WO 9612741, report that ligands which bind CD23, e.g., monoclonal antibodies, are useful in the treatment or

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prophylaxis of inflammatory, autoimmune and allergic diseases.

The usage of monoclonal antibodies to CD23, as both IgE agonists and antagonists has been reported. IgE antagonists have been reported to have potential utility in treatment of conditions or diseases wherein IgE suppression is therapeutically desirable, e.g., allergic conditions such as allergic rhinitis and conjunctivitis, atopic dermatitis and asthma. For example, Bonnefoy et al., WO 8707302 (1987), report monoclonal antibodies to human CD23, which are assertedly useful for assaying the presence of IgE receptors on cell types and as therapeutics in diseases wherein modulation of IgE is therapeutically desirable.

In part because of their potential as therapeutics and diagnostics, many groups have reported the generation of monoclonal antibodies to CD23. See, e.g., Rector et al., *Immunol.*, 55:481-488 (1985); Suemura et al., *J. Immunol.*, 137:1214-1220 (1986); Noro et al., *J. Immunol.*, 137:1258-1263 (1986); Bonnefoy et al., *J. Immunol.*, 138:2170-2178 (1987); Flores-Romo et al., *Science*, 261:1038-1046 (1993); Sherr et al.; *J. Immunol.*, 142:481-489 (1989); and Pene et al., *Proc. Natl. Acad. Sci., USA*, 85:6880-6884 (1988). Moreover, as discussed *supra*, the usage of such antibodies specifically to inhibit IgE production in systems where IgE synthesis is cytokine (IL-4) induced has also been reported. (Flores-Romo et al (*Id.*); Sherr et al. (*Id.*); Bonnefoy et al. (WO 8707302); Bonnefoy et al. (WO 8707302); Bonnefoy et al. (WO 9612741)); Bonnefoy et al., *Eur. J. Immunol* 20:139-144 (1990); Sarfati et al., *J. Immunol* 141:2195-2199 (1988) and Wakai et al.,

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Hybridoma 12:25-43 (1993). Also, Flores-Romo et al. (Id.) disclose that Fabs prepared from anti-CD23 antibodies inhibit antigen-specific induced IgE responses *in vivo* in the rat. However, notwithstanding what has been reported, the mechanism by which anti-CD23 antibodies modulate IgE expression and in particular, the manner by which they block IL-4 induced IgE production remains unclear.

It has been suggested that anti-CD23 antibodies inhibit IgE production by signaling through CD23 present on the surface of IgE secreting B cells. It has been proposed that the function of CD23, which is upregulated on IgE secreting B cells, is feedback inhibition of IgE production (Yu, et al. *Nature* 369, 753-756 (1994)). This has been theorized because mice in which the CD23 gene has been removed have increased and sustained IgE production compared to controls (Yu, et al.). In addition, it has been reported that binding to CD23 by IgE complexes or by a monoclonal antibody to anti-CD23 suppresses ongoing IgE synthesis by a lymphoblastoid cell line that constitutively secretes IgE (Sherr et al. (Id.)). It appears that this is due to down regulation of the messenger RNA for the secreted IgE heavy chain in this cell (Saxon et al., *J. Immunol.*, 147:4000-4006 (1991)). However, the exact mechanism by which IgE expression is inhibited has yet to be explained in systems in which IgE secretion is IL-4 induced.

It has also been reported that crosslinking of Fc gamma RII with surface Ig (B cell receptor) on B cells leads to down regulation of Ig expression. (D'Ambrosia et al., *Science*, 268:293-297 (1995). A similar mechanism can be proposed for B cells secreting IgE which also have cell surface CD23 and Fc gamma RII. An

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anti-human CD23 antibody bound to a cell by antigen (CD23) and also bound to Fc gamma RII through Fc interactions could transmit a signal to suppress IgE secretion through Fc gamma RII.

5 Mechanisms involved in IgE inhibition by anti-CD23 antibodies have been proposed that include blocking interactions other than the interaction between membrane CD23 and IgE. Related to this, CD23, which is a member of the C-type lectin family, has been shown to interact
10 with several other ligands such as CD21, CD11b and CD11c present on a variety of cell types including T cells and monocytes. In this context CD23 can be envisioned as a cellular adhesion molecule.

 Therefore, it has been proposed that the CD21-CD23
15 interaction may be involved in antigen presentation and subsequent IgE production. Models suggest CD21 on B cells sending an activation signal for IgE production after binding to CD23 on activated T cells present primarily in atopic individuals. (Leconant et al.,
20 *Immunol.*, 88:35-39 (1996); and Bonnefoy et al., *Int. Amer. Allergy Immunol.*, 107:40-42 (1995).) Blocking this interaction with an anti-CD23 could block induced IgE production. (Aubry et al., *Nature*, 358:505-507 (1992) and *Immunol.*, 5:944-949 (1993); Grosjean et al.
25 (1992); Bonnefoy et al., *Curr. Opin. Eur. J. Immunol.*, 24:2982-2988 (1994); Henchoz-Lecoanet et al., *Immunol.*, 88:35-39 (1996) Nambu et al., *Immunol. Lett.*, 44:163-167 (1995); Bonnefoy et al., *Int. Amer. Allergy Immunol.*, 107:40-42 (1995).) It is also possible that antigen
30 presentation is upregulated by CD23 on antigen presenting B cells binding to CD21 on T cells.

 Yet another mechanism which would potentially explain the effects of CD23 on IgE production involves

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soluble forms of CD23. It has been reported that CD23 is cleaved from the cell surface releasing several different forms of soluble CD23 or IgE binding factors. (Sarfati et al., *Immunol.*, 53:197-205 (1984).) Soluble

5 CD23 is a cytokine, with one of its reported activities being the augmentation of IL-4 induced IgE production from B cells. (Pene et al., *J. Cell Biochem.*, 39:253-269 (1989); Pene et al., *Eur. J. Immunol.*, 18:929-935 (1988); Sarfati et al., *J. Immunol.*, 141:2195-2197 (1988); Sarfati et al. (1984) (*Id.*);

10 (Saxon et al., *J. Clin. Immunol. Allergy*, 86 (3 pt 1) 333-344 (1990). Also, certain forms of soluble CD23 have been reported to inhibit IgE production (Sarfati et al., *Immunol.*, 76:662-667 (1992)). Accordingly,

15 anti-CD23 antibodies potentially may block IgE production by 1) inhibiting the IgE augmenting effects of soluble CD23 and/or 2) blocking the proteolytic release of soluble CD23 from the cell surface.

Thus, based on the foregoing, it is clear that

20 there is significant complexity and uncertainty in the art with respect to the functions of more specifically CD23 and effects on IgE production, and further with respect to the means by which ligands specific thereto affect IgE production.

25 **Objects of the Invention**

Thus, it is an object of the invention to produce novel ligands (antibodies) specific to CD23 and to use such antibodies to elucidate the mechanism by which anti-CD23 antibodies modulate IgE expression.

30 It is another object of the invention to produce novel ligands (antibodies) which bind CD23, in

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particular human CD23, having improved ability to inhibit induced IgE expression.

It is a more specific object of the invention to produce anti-human CD23 antibodies containing human
5 gamma-1 or human gamma-3 constant domains.

It is another object of the invention to produce multivalent anti-human CD23 antibodies which may be more effective by virtue of their enhanced potential for cross linking CD23 and Fc receptors.

10 It is another object of the invention to provide pharmaceutical compositions containing anti-human CD23 monoclonal antibodies comprising human gamma-1 or gamma-3 constant domains which are capable of inhibiting induced IgE production.

15 It is another object of the invention to use an anti-human CD23 monoclonal antibody comprising a human gamma-1 or human gamma-3 constant domain for treatment or prophylaxis of disease conditions wherein inhibition of induced IgE production is therapeutically desirable.

20 More specifically, it is an object of the invention to treat or prevent allergic conditions, autoimmune diseases and inflammatory diseases using an anti-human CD23 monoclonal antibody comprising either a human gamma-1 or human gamma-3 constant domain.

25 **Brief Description of the Figures**

Figure 1 compares the *in vitro* IgE inhibitory activity of a murine anti-human CD23 monoclonal antibody (MHM6), to five primate anti-human CD23 monoclonal antibodies (5E8, 6G5, 2C8, B3B11, and 3G12);

30 Figure 2 shows that primate monoclonal antibodies 5E8 and 6G5 bind an epitope on human CD23 that is distinct from commercially available murine anti-human CD23 monoclonal antibody MHM6 (middle panel, figure 2)

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and compete with each other (lower panel, figure 2).
Primate anti-human CD23 monoclonal antibodies 2C8 and,
B3B11 compete with MHM6 top panel, figure 2).

Figure 3 compares the *in vitro* IgE inhibitory
activity of a particular primate anti-human CD23
monoclonal antibody 5E8 to four different PRIMATIZED®
versions of said primate monoclonal antibody, the
sequences of which are described below.

p5E8G4P- This PRIMATIZED® antibody contains the
following sequences:
Human kappa light chain constant region
and a human gamma 4 constant region which
contains a P mutation (Angal et al., *Mol.*
Immunol., 30:105-108 (1993));

p5E8G4PN- This PRIMATIZED® antibody contains the
human kappa light chain constant region
and a human gamma 4 constant region
having a P mutation (Angal et al. *Mol.*
Immunol., 30:105-108 (1993)). This
antibody also contains a mutation in the
heavy chain variable region which changes
an asparagine residue (potential
carbohydrate attachment site) to a
lysine;

p5E8G1 This PRIMATIZED® antibody contains the
human kappa light chain constant region
and a human gamma 1 constant region;

p5E8G1N- This PRIMATIZED® antibody contains the
human kappa light chain constant region
and human gamma 1 constant region. This
antibody also contains a mutation in the
heavy chain variable region which changes

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an asparagine residue (carbohydrate attachment site) to a lysine;

Figure 4 contains a table which compares the apparent Kd in nM of the antibodies identified in Figure 3 and summarizes their IgE suppressive activity.

Figure 5 compares the *in vitro* IgE inhibitory activity of a particular primate anti-human CD23 monoclonal antibody, 6G5, to two different PRIMATIZED® versions of 6G5 which are described below:

- | | | |
|----|---------|--|
| 10 | p6G5G1 | This PRIMATIZED® antibody contains the human lambda light chain constant region and the human gamma 1 constant region; |
| 15 | p6G5G4P | This PRIMATIZED® antibody contains the human lamda light chain constant region and the human gamma 4 constant region with a P mutation (Angal et al., <i>Mol. Immunol.</i> , 30:105-108 (1993)); |

Figure 6 compares the *in vitro* IgE inhibitory activity of primate anti-human CD23 monoclonal antibody 2C8 to F(ab')₂ derived from 2C8;

Figure 7 shows that the F(ab')₂ derived from 2C8 antagonizes the suppression of *in vitro* IgE activity of primate anti-human CD23 monoclonal antibody 2C8.

Figure 8 shows the *in vivo* IgE inhibitory activity of a particular primate anti-human CD23 monoclonal antibody, 5E8, in a SCID animal model;

Figure 9 compares the *in vivo* inhibitory activity of primate anti-human 6G5 and a PRIMATIZED® version thereof p6GSG4P.

Figure 10 shows the *in vivo* IgE inhibitory activity of the primate anti-human CD23 monoclonal antibody 6G5 and a PRIMATIZED® version thereof, p6G5G1.

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Definition of Terms Used in This Application

Chimeric Antibody:

5 A recombinant antibody containing regions from two different antibodies, usually different species antibodies, most typically rodent variable sequences and human constant domain sequences.

Anti-Human CD23 Gamma 1 Antibody

10 An antibody that specifically binds human CD23 which contains a human gamma 1 constant region or fragment or modification thereof which inhibits induced IgE production. This includes, in particular, antibodies containing rodent or primate variable domains or antigen binding portions, humanized, PRIMATIZED®, and
15 human anti-human CD23 monoclonal antibodies which comprise a human gamma 1 constant domain, fragment, or modification thereof, which inhibit induced IgE production *in vitro*.

Anti-Human CD23 Gamma 3 Antibody

20 An antibody that specifically binds human CD23 which contains a human gamma 3 constant region or fragment or modification thereof which inhibits induced IgE production. This includes, in particular, antibodies containing rodent or primate variable domains or antigen binding portions, humanized, PRIMATIZED®, and
25 human anti-human CD23 monoclonal antibodies which comprise a human gamma 3 constant domain, fragment, or modification thereof, which inhibit induced IgE production *in vitro*.

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Modifications of Antibody Constant Domains

Antibodies according to the present invention containing mutations, substitutions or deletions of the constant region, that may create a desired change in the level of efficiency, i.e. in FcR binding, without changing the basic effector functions mediated by the constant region.

PRIMATIZED® Antibody

A recombinant antibody containing primate variable sequences or antigen binding portions, and human constant domain sequences.

Humanized Antibody:

A recombinant antibody containing a non-human variable region or antigen binding portion which has been modified to more closely mimic a human antibody variable region and thereby eliminate or minimize potential immunogenicity if administered to humans without sacrificing the specificity or affinity of the immunoglobulin. There are several known methods of humanization, including, "veneering" which comprises select modification of surface residues, framework replacement, (CDR grafting) and molecular modeling.

Gamma 1 Constant Domain:

A particular type of constant domain sequence which confers upon an antibody specific effector activities. In the present application, gamma 1 constant domain refers to a human gamma 1 constant domain, fragment or modification thereof, which retains gamma 1 effector functions in combination with anti-CD23 variable domain sequences or antigen binding portions. Modifications include human gamma-1 constant domains which comprise

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the deletion, substitution or addition of one or more amino acid residues. This effector function is manifested by the ability of an antibody containing such a constant domain to inhibit induced IgE production.

5 Gamma 3 Constant Domain:

 A particular type of constant domain sequence which confers upon an antibody specific effector activities. In the present application, gamma 3 constant domain refers to a human gamma 3 constant domain, fragment or
10 modification thereof, which retains gamma 3 effector functions in combination with anti-CD23 variable domain sequences or antigen binding portions. Modifications include human gamma-3 constant domains which comprise the deletion, substitution or addition of one or more
15 amino acid residues. This effector function is manifested by the ability of an antibody containing such a constant domain to inhibit induced IgE production.

CD23:

 This refers to the low affinity receptor for IgE,
20 FcεRII/CD23.

Anti-CD23 Antibody:

 An antibody that specifically binds CD23, preferably human CD23.

Detailed Description of the Invention

25 As discussed *supra*, while many groups have previously reported the production of anti-CD23 antibodies and the use thereof as antagonists and agonists for modulating IgE production, the exact mechanism by which such antibodies modulate IgE
30 expression in systems where IL-4 induces IgE production

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remains unclear. Thus, it would be beneficial if the means by which such antibodies modulate IgE expression were elucidated, or at least better explained, as such information would be potentially useful in designing therapeutics for treatment of diseases wherein modulation of IgE production is therapeutically desirable. In particular, it would be beneficial if improved antibodies specific to CD23 were obtained having improved capacity to inhibit induced IgE production, as enhanced IgE levels are believed to be involved in numerous disease processes, e.g., allergic conditions, inflammatory conditions and autoimmune diseases. Such diseases include by way of example, atopic dermatitis, eczema, allergic rhinitis and conjunctivitis, Job's syndrome, and asthma.

Toward that end, the present inventors have surprisingly discovered that anti-human CD23 monoclonal antibodies which contain human gamma-1 constant domains inhibit IgE production in systems where IgE production is induced by IL-4 significantly better than CD23 monoclonal antibodies of other effector types, e.g., those comprising human gamma-4 constant domains or CD23 monoclonal antibodies or antibody fragments lacking effector functions altogether. Because human gamma-3 constant domains have been shown to mediate the same effector functions as human gamma-1, anti-human CD23 monoclonal antibodies containing human gamma-3 constant domains are also included.

There are currently five defined effector functions for the IgG (gamma) class of antibodies. Two of these functions, complement activation and FcγRN interaction, are not found in the in vitro assays described in the present invention, and are therefore not likely to be involved in the molecular mechanism. Three other FcγR

receptors have been identified which interact with the IgG class of antibodies: FcγRI, FcγRII (of which there are at least six different proteins) and FcγRIII (with at least two different proteins). All three of these
5 receptors interact with both IgG1 and IgG3.

FcγRI is the only one of the three having an appreciable affinity for IgG. It binds both monomeric gamma-1 and gamma-3 with a K_a of about $5 \times 10^8 \text{ M}^{-1}$. However, its affinity for human gamma-4 is about 10-fold
10 less, and it does not bind human gamma-2 at all (Fries et al., 1982, J. Immunol. 129: 1041-1049; Kurlander and Batker, 1982, J. Clin. Invest. 69: 1-8; Woof, 1986, G. Mol. Immunol. 21: 523-527; see also Burton and Woof, 1992, Human Antibody Effector Function, Adv. Immunol.
15 51: 1-84).

Although the affinity of human FcγRII and FcγRIII for human IgG is generally very low ($K_a < 10^7 \text{ M}^{-1}$), affinity for human IgG1 and human IgG3 increases significantly ($K_a \approx 2 \text{ to } 5 \times 10^7 \text{ M}^{-1}$) when they are bound to antigen
20 (Karas et al., 1982, Blood 60: 1277-1282). The affinity of human FcγRII for human IgG2 bound to antigen has given conflicting results. Human FcγRIII does not bind to human IgG2. Human FcγRII and human FcγRIII do not bind to human IgG4 (Van de Winkel and Anderson, 1991, J.
25 Leuk. Biol. 49: 511-524; Huizinga et al., 1989, J. Immunol. 142: 2359-2364).

While Fc mediated effector functions are sometimes significant to the therapeutic activity of antibodies, this discovery was surprising in the case of anti-CD23
30 antibodies because the role of effector function in the IgE inhibitory activity of anti-CD23 antibodies had not been previously reported. In fact, previous evidence had suggested that antibody effector function was not significant to the ability of anti-CD23 antibodies to

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inhibit induced IgE production. For example, Flores-Romo et al., *Science*, 261:1038-1041 (1993) had reported that Fabs prepared from a polyclonal anti-CD23 antibody inhibited an *in vivo* induced IgE

5 antigen-specific response.

The discovery that effector functions mediated through the constant region of the anti-CD23 antibodies are apparently involved was made after the present inventors isolated various primate antibodies specific to CD23 having anti-IgE inhibiting activity and compared these antibodies to PRIMATIZED® versions with respect to their ability to inhibit IL-4 induced IgE production *in vitro* and *in vivo*. Antibodies constructed with a human gamma-4 constant region failed to inhibit IgE antigen-specific responses *in vitro*, whereas antibodies containing a human gamma-1 constant region were successful.

Because one (or more) of the three classes of FcγR receptors, FcγRI, FcγRII or FcγRIII, is likely involved in the specific effector function mediated through the gamma-1 domain, and because these classes of receptors also bind to antibodies containing gamma-3 domains, it is logical that anti-CD23 antibodies containing gamma-3 domains will also be successful at inhibiting IgE-antigen specific-response *in vitro*.

More specifically, and as described in greater detail *infra*, five primate monoclonal antibodies which specifically bound both cellular and soluble CD23 were isolated from an Old World monkey (macaque) according to the methodology which is disclosed in commonly assigned Application Serial No. 08/379,072 (now allowed), which application is incorporated by reference in its entirety herein. This application described in detail a means

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for producing monoclonal antibodies to desired antigens, desirable human antigens, in Old World monkeys and their advantages in relation to antibodies of other species as therapeutics, for example reduced or potentially lack of immunogenicity in humans because of the phylogenetic closeness of humans and Old World monkeys. In fact, because of the phylogenetic closeness of these species, it is difficult to distinguish Old World monkey immunoglobulins from human immunoglobulins by sequence comparison.

Four of these five primate monoclonal anti-human CD23 antibodies were demonstrated to be capable of inhibiting IL-4 induced IgE production in an *in vitro* B cell assay described in detail *infra* and the most potent was also shown to inhibit IL-4 induced IgE in a SCID mouse animal model (also described in detail *infra*). Based on this IgE inhibitory activity, and expected low immunogenicity in humans, such antibodies are potentially suitable as therapeutics for treating diseases wherein inhibition of IgE production is therapeutically desirable.

However, in order to further reduce immunogenicity, it was elected to PRIMATIZE® two primate monoclonal antibodies (a type of chimerization of antibodies) according to the methodology which is also described in U.S. Serial No. 08/379,072 (now allowed), incorporated by reference herein. PRIMATIZATION® essentially refers to the production of recombinant antibodies developed by IDEC Pharmaceuticals Corporation which comprise primate variable regions and human constant regions. Primatization of the two primate anti-human CD23 monoclonal (5E8 and 6G5) antibodies having potent IgE inhibiting activity was effected in order to eliminate

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any potential immunogenicity attributable to the primate constant domains in humans.

Again, because of the inventors' initial expectation from published literature that Fc effector function was not necessary for induced IgE inhibition, human gamma 4 versions of these particular antibodies were initially produced. However, quite surprisingly, it was found that the gamma-4 versions produced from both of these primate monoclonal antibodies were ineffective, i.e., they required significantly higher concentrations of PRIMATIZED® gamma 4 antibody than the primate antibody to inhibit IL-4 induced IgE production in *in vitro* assays.

Moreover, even more surprising was the discovery that when the same two primate antibodies were then converted to human gamma-1 versions (by substitution of the primate constant domains with human gamma-1 constant domains), that these gamma-1 antibodies very effectively inhibited induced IgE production *in vitro*. Thus, our results suggested that Fc effector function is apparently significant to the ability of anti-human CD23 antibodies to inhibit induced IgE production. This hypothesis was confirmed when a third primate anti-human CD23 monoclonal, i.e., the 2C8 antibody, which was shown by us to inhibit IgE production *in vitro*, was converted to a F(ab')₂, which was found to be substantially incapable of inhibiting induced IgE production *in vitro*. In fact, this F(ab')₂ was found to antagonize the suppressive effects on induced IgE blocking activity of the primate anti-human CD23 monoclonal antibody 2C8.

In addition, it was found that removing a glycosylation site in the heavy chain variable region of one of the antibodies (5E8) had no effect on binding of the antibody to CD23 (as evidenced by obtained Kd

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values), or on induced IgE inhibition. Thus, the differences in IgE inhibition were shown to apparently not involve glycosylation differences.

5 The PRIMATIZED® gamma 1 version of primate 6G5 was found to inhibit induced IgE expression in SCID mice while the same concentration of either the primate 6G5 or the PRIMATIZED® p6G5F4p did not inhibit induced IgE expression. Therefore, an antibody containing human gamma-1 constant domains was found to be even more
10 effective in an *in vivo* animal model than the primate monoclonal antibody. Furthermore, the inventors anticipate that anti-CD23 antibodies containing human gamma-3 constant domains will be just as effective as those having gamma-1 constant domains, because gamma-1
15 and gamma-3 constant domains have affinity for the same classes of Fc receptors.

Accordingly, based on these results, it has been surprisingly discovered that an active Fc region, in particular that of human gamma-1 or human gamma-3, is
20 significantly involved in the mechanism of IL-4 induced IgE inhibition by anti-human CD23 monoclonal antibodies. This discovery is quite unexpected especially based on earlier reports that Fabs derived from polyclonal anti-CD23 antibodies were capable of inhibiting induced
25 IgE production, and also based on the various theories as to how CD23 affects induced IgE expression.

Accordingly, the present invention relates to anti-human CD23 antibodies containing human gamma-1 or gamma-3 constant domains and their use as therapeutics
30 based on their ability to effectively inhibit IgE expression.

The skilled artisan can prepare anti-human CD23 antibodies containing either human gamma-1 or gamma-3 constant domains by methods which are well known in the

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art for the manufacture of chimeric antibodies.

Essentially, such methods comprise producing anti-human CD23 antibodies in a desired host or *in vitro*, cloning a hybridoma or cell line which produces an anti-human CD23 monoclonal antibody exhibiting desirable characteristics, e.g., adequate CD23 binding affinity, cloning the nucleic acid sequences which encode such antibody from said hybridoma or cell line, e.g. by polymerase chain reaction using suitable primers, isolating the variable domains contained therein, recombining such variable domains with human gamma-1 or gamma-3 constant domains and the appropriate human light chain constant domain, and expressing the resultant nucleic acid sequence encoding a chimeric anti-human CD23 gamma-1 or gamma-3 immunoglobulin in a suitable expression system. Preferably, the anti-human CD23 antibodies of the invention will have apparent CD23 binding affinities ranging from 0.1 nM to 1000 nM, more preferably at least 50 nM, and most preferably at least 5 nM.

Host cells suitable for expression of recombinant immunoglobulins are well known in the art. For example, recombinant antibodies may be expressed in Chinese hamster ovary (CHO) cells, DG44 or DUXB11; or CHO cells CHO K-1; mouse myeloma cells SP2/0 or X63-Ag8.653 or NSO; rat myeloma cells YB2/0; baby hamster kidney cells, BHK; human embryonic kidney line, 293; monkey kidney cells, CV1; human lung fibroblasts, WI38; human cervical carcinoma cells, HELA; insect cells, plant cells, yeast or in bacteria. Further, vectors suitable for expression of immunoglobulins are also well known in the art and are commercially available.

A particularly preferred vector system is the translationally impaired vector system disclosed in U.S. Serial No. 08/147,696 (now allowed), which comprises a

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translationally impaired dominant selectable marker (neo) containing an intron into which a desired heterologous DNA is inserted. This vector system has been found to provide for very high yields of recombinant proteins, e.g., immunoglobulins. However, the subject anti-CD23 antibodies may be produced in any vector system which is suitable for expression of functional immunoglobulins.

Also, the present invention embraces human monoclonal antibodies of the gamma-1 or gamma-3 types which are specific to human CD23. Methods for isolation of human monoclonal antibodies are also well known in the art and include *in vitro* methods, e.g., *in vitro* immunization of human B cells in tissue culture, and *in vivo* methods, e.g. synthesis of human monoclonal antibodies in SCID mice. A preferred means of producing human monoclonal antibodies in SCID mice which combines *in vitro* priming of human spleen cells which are then introduced into SCID mice is disclosed in U.S. Serial No. 08/488,376 (incorporated by reference in its entirety herein). This method is advantageous as it provides for the reproducible recovery of monoclonal antibodies having high affinity against a desired antigen, e.g., a human antigen.

Also, the present invention embraces human monoclonal antibodies which compete with the primate anti-human CD23 monoclonal antibodies 5E8 and 6G5 for binding to CD23.

EXAMPLE 1

Production of Primate Anti-CD23 Antibodies

Five primate monoclonal antibodies specific to CD23 were isolated from macaques substantially according to

the methodology disclosed in Serial No. 08/379,072, which has been incorporated by reference herein. The exact techniques utilized are described in detail below.

Methodology for Isolation and Characterization
of Anti-Human CD23 Monoclonal Antibodies

Purification of the Immunogen sCD23 from 8866 cells

During purification, soluble CD23(sCD23) was quantified by a three-step ELISA using a murine anti-CD23 antibody (Binding Site; catalog # MC112) as a capture. The antigen was partially purified from cultures of 8866 cells maintained in suspension bioreactors using RPMI 1640 (JRH Biosciences; catalog # 56-509) supplemented with 10% fetal bovine serum (JRH Biosciences) and 4 mM glutamine (JRH Biosciences; catalog # 90114) at 37°C. Carbon dioxide was used to maintain pH 7.1. After removing cells by 0.45 µm filtration, phenylmethyl sulfonyl fluoride (final concentration 0.2 mM, Sigman Chemical Co.; catalog # P-7626) and ethylenediaminetetraacetic acid (final concentration 3 mM, Sigma Chemical Co.; catalog # EDS) were added to the supernate and the solution stored at 2-8°C. The cell-free supernate was concentrated approximately 15 to 20-fold using a hollow-fiber ultrafiltration cartridge (A/T Technology; catalog # UFP-10-C-9A; 10,000 d MWCO) or tangential flow ultrafiltration cartridge (Filtron Corporation; 10,000 d MWCO) at ambient temperature. The concentrated supernate was sterile filtered and stored at -70°C. Thawed concentrates were de-lipidated by adding SM-2 BioBeads (BioRad Industries; catalog # 152-3920) at 5 g/L and stirring overnight at 2-8°C. The resin was removed by filtration and the solution stored at 2-8°C. For some preparations of sCD23, concentrates were

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fractionated using ammonium sulfate (35-70% (w/v); Fisher; catalog # A702-3) before or after de-lipidation.

5 The de-lipidated solution was subsequently purified using affinity chromatography at 2-8°C. The affinity matrix was prepared by covalently linking a murine anti-CD23 monoclonal antibody (BU38) to Sepharose using CNBr-activated Sepharose 4B (Sigma Chemical Co.; catalog # C-9142). The BU38 antibody was purified to >90% homogeneity from ascites (Binding Site; catalog # CUS830) using Protein A chromatography. The de-lipidated solution was applied to the affinity column (1.5 x 5 cm) equilibrated with 1XPBS (Gibco BRL; catalog # 70013-0.32), pH 7.2 and the column washed with 1XPBS, pH 7.2, containing 0.05% NP40 (Sigma Chemical Co.) to remove non-bound protein. Soluble CD23 was eluted using 3.5 M MgCl₂ (Fisher; catalog # M33-500). Fractions containing sCD23 were combined and dialyzed (Baxter Spectra/Por; catalog # D1615-1) against 1XPBS, pH 7.2 at 2-8°C. After dialysis, the protein solution was concentrated by centrifugation using Centriprep 10 spin filters (Amicon Corporation; MWCO 10,000 d) and preparations stored at -70°C. The purity of sCD23 was estimated to be >70% using SDS-PAGE analysis (4-20% pre-cast gels, Novex Corporation) and Coomassie staining.

25 **Immunization of Primates and Isolation of Immune Cells**

Cynomolgus monkeys (White Sands Research Center, Alamogordo, New Mexico) were immunized with soluble CD23 which had been purified from the supernatant of human RPMI 8866 cells (B cell lymphoma, Hassner and Saxon, J. Immunol., 132:2844 (1984)). Each monkey was immunized every third week with 200 µg soluble CD23 in 500 µl PBS mixed with 167 µl Temuritide (adjuvant peptide) (Sigma, St. Louis, MO, Catalog # A-9519) and 333 µl 3X PROVAX®

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(IDEC Pharmaceuticals Corporation). Immunization was effected intradermally, intraperitoneally, intramuscularly and subcutaneously. The titer of anti-CD23 antibodies in the serum of the monkeys was measured by ELISA on 8866 cells and compared to a pre-bleed from the same monkeys.

Monkey PRO 978, with a serum titer of fifty thousand was sacrificed, and the spleen and lymph nodes were surgically removed, and shipped on ice to IDEC pharmaceuticals, submerged in sterile RPMI-1640 (Gibco BRL, Gaithersburg, MD, Catalog # 21870-050) supplemented with 10% fetal calf serum, 2 mM L-glutamine, 2 mM sodium pyruvate and 50 μ g/ml gentamicin. Immediately upon arrival the spleen was homogenized by squeezing it through a wire mesh with a glass pistil. Red blood cells were lysed in an ammonium chloride based hypotonic buffer and the remaining lymphocytes collected and washed in RPMI-1640 at least three times. Lymph nodes were homogenized similarly into a single cell suspension, collected and washed at least three times in RPMI-1640.

Production of Hybridomas

After the last wash, the cells were counted, and the primate cells obtained above were then somatically fused to the mouse-human heterohybridoma cell line H6K6/B5 (Carroll et al., *J. Immunol. Methods*, 89:61 (1986)) using standard techniques (Boerner et al., *J. Immunol.*, 147:86) (1991)) and plated into 96 well dishes (175 dishes or 14,700 wells for the spleen, and 17 dishes or 1386 wells for the lymph nodes) at 300,000 cells per well.

This procedure involved the mixing of lymphocytes and the above-identified fusion partner, at a 2:1 ratio,

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which cells were slowly resuspended into 50% PEG 1500 (Sigma, Catalog # P5402) for 1 minute. These cells were then allowed to rest for 1 minute and then slowly further resuspended in excess RPMI-1640. Afterward, the cells were again allowed to rest, this time for 15 minutes before a light spin at 250 x g. The cells were then resuspended in RMPI-1640 growth media, which was supplemented with 20% Fetal Calf Serum, 2 mM L-Glutamine, Sodium Pyruvate, Non-Essential Amino Acids and 50 µg/ml Gentamicin, containing 100 µM Hypoxanthine, 16 µM Thymidine (BoehringerMannheim, Germany, # 623091) and 5.8 µM Azaserine (Sigma, Catalog # A 1164) (HTA). HTA is a selection agent which provides the survival of successfully fused cells (primate lymphocyte fused with heterohybridoma fusion partner).

Approximately 65% of the wells showed growth (10,500 wells). These wells were then screened for the presence of anti-human CD23 antibody by a three step cell ELISA.

ELISA Procedure

The first step of the ELISA comprised the transferral of fifty microliters of supernatant from each well to ninety-six well plates which had previously been coated with 10⁵ 8866 cells (CD23 positive cell line) per well. These plates were made by first coating the plates with 50 µl of aqueous solution containing twenty µg/ml Poly L-Lysine (Sigma Catalog # P1399, MW 150,000 - 300,000) for thirty minutes at room temperature. The remaining solution was removed ("flicked out") and the plates left to dry. Once dry, fifty µl of 8866 cells in PBS were transferred and spun at 600 g for five minutes. The 8866 cells were covalently bound to the plate by adding fifty µl 0.5% glutaraldehyde (Sigma Catalog #

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G6257) in phosphate buffered saline (PBS) for 15 minutes. The glutaraldehyde was removed (flicked out) and the plates blocked with one hundred fifty μ l 100 mM glycine (Sigma Catalog # G-2879) in 0.1% BSA - PBS.

5 After the addition of supernatants, the plates were incubated at 37°C for one to two hours and washed seven to nine times with tap water, and a goat anti-human IgG antibody coupled to horse radish peroxidase (HRPO) (Southern Biotech, Birmingham, Alabama, Catalog #

10 2040-05) diluted 1:2000 into 1% dry skimmed milk (Vons) in PBS - 0.05% Tween 20 (Sigma, Catalog # P1379) was added. The plates were incubated for forty-five minutes at 37°C, and again washed seven to nine times in tap water. The presence of the HRPO was detected by a color

15 development after the addition of a TMB reagent (Kirkegaard & Perry, Gaithersburg, MD, Catalog # 50-76-02 and 50-65-02), 100 μ l/well. The reaction was stopped by adding twenty-five μ l 4N H₂SO₄. Optical density (OD) was measured at 470 nM on a

20 spectrophotometer (Titertek Multiscan). The OD values greater than two times the background were scored as positive.

The second step in the ELISA was effected to confirm that the supernatants which had been scored

25 positive in the first ELISA reacted to CD23 and not to some irrelevant antigen. This was effected by testing the supernatants on SupT1 cells (ERC BioServices Corporation, Rockville, MD, Catalog # 100), a CD23 negative human cell line, using the same ELISA

30 procedure. Supernatants that scored similarly in both tests were discarded. These results indicated that fifty-six of the 10,500 wells with growth showed the presence of a primate monoclonal antibody that bound to

8866 cells in two separate screenings at different times and did not bind to SupT1 cells.

5 The third step of the ELISA was conducted to determine whether the supernatants identified according to the first two ELISA steps, reacted with soluble CD23. In this third ELISA, 96 well plates were coated at 4°C overnight with 2 µg/ml BG-6 (Biosource International, Camarillo, CA, Catalog # CT-CD23-CF), a mouse monoclonal antibody that binds to soluble CD23 but does not block
10 CD23-IgE binding, contained in a 50 mM bicarbonate buffer, pH 9.3. After removing the coating buffer, fifty µl of semi-purified soluble CD23 at a predetermined dilution in PBS were added to the plate and incubated for two hours at room temperature. After
15 washing the plate with tap water seven to nine times, 50 µl supernatants from selected wells were added. After washing the plate in tap water seven to nine times, 50 µl rabbit anti-human IgG (mouse adsorbed)-HRPO (Southern Biotech, Catalog # 6145-05) diluted 1:4000 in 1% dry skimmed milk in PBS with 0.05% Tween 20 were added,
20 incubated for two hours at 37°C, washed seven to nine times in tap water and developed with TMB as described above. Wells with OD's greater than two times the background were again scored as positive.

25 Twenty-one of the fifty-six wells that showed binding to 8866 cells also bound to sCD23 in the ELISA. These wells were expanded and subcloned at least twice by plating out cells at one cell per three wells. After approximately three months, five stable hybridomas
30 producing primate monoclonal antibodies to CD23 were obtained.

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Antibody Purification by Protein A Methods

Essentially, antibodies are purified by centrifuga-
tion of the culture supernatant to remove cells and de-
bris. The resultant centrifuged samples are then
5 filtered through a 0.2 μ m filter. A protein A sepharose
Fast flow column is then prepared and equilibrated using
PBS (pH 7.4). The supernatant is then loaded on the
column at an appropriate flow rate (2 ml/min). After
loading, the wash column is washed with 10 column volume
10 of PBS (pH 7.4). The antibody is then eluted from the
column with elution buffer (0.2 M acetic acid, 0.1 M
glycine pH 3.5) at 1 ml/min flow rate. One milliliter
fractions/tube (2.0 M Tris-Hcl pH 10.0) including 100 μ l
of Tris, are then collected. Afterward,
15 spectrophotometer readings are taken at 280 nm. The
resultant fractions with high absorbance at 280 nm
containing the antibody are then collected and dialyzed
against PBS overnight. The product is then sterilized
by filtration through 0.22 μ m membrane and stored at -
20°C.
20

Four of these five primate anti-human CD23
monoclonal antibodies (1H6, 2C8, 5E8 and 6G5) were
demonstrated to inhibit IgE production in an *in vitro*
assay which measures IgE production by
25 IL4-hydrocortisone induced peripheral blood mononuclear
cell (PBMC) cultures. These results are shown in Figure
1. The assay conditions are described below. The fifth
primate monoclonal anti-human CD23 antibody B3B11 was
inactive in this assay.

30 **IL-4 Stimulated IgE Production**
by Peripheral Blood Mononuclear Cells

As discussed supra, the subject primate antibodies
and PRIMATIZED® forms thereof were assessed for their

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[illegible]

5

10

anti-CD23 Mabs:

primate Mabs (no preservatives)

15

HB101 supplement: (Irvine Scientific Catalog #
T151)

20

hydrocortisone: (Sigma Catalog # H-0888)

25

Histopaque®: (Sigma Catalog # H-8889)

1% FBS in HBSS

30

Bath Clear Microbicide (Fisher, Catalog #

13-641-334) in DPBS

puromycin solution: 40 μ g/ml in HB101 growth medium

hydrocortisone solution: 0.1 M solution in DMSO
anti-CD23 murine Mabs were extensively dialyzed to
remove preservatives

HB101 growth medium

Buffy coat cells 1:4 are diluted using HBSS at room temperature. These cells are derived from whole blood after an overnight incubation at room temperature to resolve and separate the plasma components, clotted platelets and fibrin, and buffy coat cells.

Thirty microliters of diluted buffy coat are then overlaid onto fifteen microliters of Histopaque in fifty ml conical tubes. These tubes are then centrifuged for twenty minutes at 1700 rpm at room temperature without brakes (IEC 216 swinging bucket rotor). The white PBMC layer is then collected using a sterile pipette, taking care not to disturb the other layers. The PBMCs (peripheral blood mononuclear cells) are the buffy coat cells which have been sedimented by centrifugation partially through a HISTOPAQUE® density gradient to form a distinctly visible white layer of cells. These cells are collected with a pipette, rinsed

with HBSS, and then counted using a hemocytometer. Typically, 300 to 600 million PBMCs can be recovered from a single 450 ml buffy coat package.

5 The collected PBMCs are then washed three times in 1% FBS/HBSS. The washed cells are collected by centrifugation for seven minutes at 1300 rpm at 7°C.

10 The number of cells collected is then determined using a hemocytometer. The cell concentration is adjusted to about three million cells per milliliter of HB101 growth medium.

15 Approximately about 1.5 million cells (0.5 ml) are then added to each well of a 48 well plate. In general, five replicate samples are prepared for each experiment. The perimeter wells of each plate are not used for cell samples. Accordingly, these wells are filled, e.g., using 0.5 ml of 0.05% BathClear/DPBS.

20 0.5 ml HB101 growth medium containing desired amounts of IL-4 and Mab is then added to the wells. The IL-4 used is recombinant DNA-generated human interleukin 4. The Mab used in the assay is a murine, primate or PRIMATIZED® antibody. Typically, IL-4 is added at a final concentrate of 100 U/ml and Mab is added at a final concentrate ranging from 0.01 to 3 µg/ml.

25 The cells are then incubated for nine to eleven days at 37°C in a moist incubator set at 5% CO₂. After incubation, the supernatant fluids are collected and the IgE content is measured.

IgE ELISA

30 The following list identifies materials and solutions used in the IgE ELISAs.

Materials and Solutions Needed for IgE ELISA
sulfuric acid, 4 M

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coating buffer: 10 mM sodium bicarbonate buffer, pH 9.6
concentrated phosphate buffered saline (10x PBS) stock
solution:

	NaH ₂ PO ₄	26.6 gm
5	Na ₂ HPO ₄	289 gm
	NaCl	1064 gm
	distilled H ₂ O	10 L

blocking buffer: 10% FBS/PBS

dilution buffer: 1% BSA/0.05% Tween 20/PBS

10 washing buffer: 0.05% Tween 20/PBS

goat anti-human IgE (epsilon chain-specific), unlabeled:
(Tago Catalog # 4104)

human IgE standard: (The Binding Site Catalog # BP094)

15 goat anti-human IgE, HRP-labeled: (Tago Catalog # AHI
0504)

TMB peroxidase substrate: (KPL Catalog # 50-76-02)

peroxidase solution B: (KPL Catalog # 50-65-02)

working substrate solution: mix substrate and Solution
B at 1:1 ratio

20 Immulon II microtiter plates (Dynatech Labs Catalog #
011-010-3455)

IgE ELISA Procedure

Each well of a microtiter plate is coated using 100
μl of a coating buffer containing 2 μg/ml goat anti-
25 human IgE.

The coated plate is then incubated overnight at
4°C.

After incubation, each well in the plate is then
washed three times with 200 μl of Tween 20/PBS. After
30 washing, the non-specific binding sites are blocked with
200 μl blocking buffer/well for 1 hour at 37°C.

One hundred μl of samples or standards are then
added to each well; which wells are then incubated over-

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night at 4°C. After incubation, the samples are tested with or without dilution. A standard concentration curve is prepared for each plate using several dilutions of IgE ranging from 0.1 to 50 ng/ml.

- 5 After overnight incubation, each plate is washed five times with Tween 20/PBS.

One hundred μ l of horseradish peroxidase (HRP) labeled goat anti-human IgE diluted 1:10,000 in dilution buffer is then added to each drained well. The plate is
10 then incubated for 4 hours at 37°C.

The plates are then washed 5 times with Tween 20/PBS and 3 times with water.

One hundred μ l of 3,3',5,5'-tetramethylbenzidine working substrate solution is then added to each well.
15 The plate is then incubated for twenty-five minutes in the dark at room temperature. After incubation the developing reaction is stopped by the addition of fifty μ l of 4 M sulfuric acid.

The absorbency is then read concurrently at 450 and
20 540 nm. The 540 nm absorbency values are subtracted as background.

Assay for Kd measurement of primate monoclonal anti-human CD23 antibodies

Scatchard Analysis Procedure

25 1. **Radiolabeling Procedure**

IODO-BEADS are washed with 100mM Phosphate Buffer, pH 7.4 twice using 1 mL of buffer per 2 beads. The beads are then dried on filter paper.

The two beads are then added to 100 μ l 125 I
30 solution, containing about 1mCi of I, diluted with 200 μ l of the phosphate buffer, and left at room temperature for 5 minutes.

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The antibody (50 μ gs) is added to the preloaded beads. The reaction time for maximal incorporation of radioactivity is 6 minutes.

5 The reaction is stopped by removing the radiolabeled antibody from the reaction vessel.

10 Gel filtration is then performed to remove excess 125 I or unincorporated 125 I from the radiolabeled antibody solution. This is effected by passing the radiolabeled antibody over a column made up of 1.5mL Sephadex-G25, 1.5 mL DEAE Sephadex-A25 and 0.5mL Amberlite. The radiolabeled antibody is eluted off in a total volume of 5 mL at a concentration of about 10 μ g/mL. (Elution Buffer: 1XPBS containing 10% Gelatin, 2% Sodium Azide and 1% BSA).

15 2. Optimization Assay (Direct Binding Study)

The specific activity of the 10 μ g/mL radiolabeled solution is determined by taking a 1 μ l sample and running the sample on a gamma counter.

Example:

- 20 • $1 \times 10^5 \text{ cpm}/\mu\text{l} \times 1000 \mu\text{l}/10\mu\text{g antibody}$
- $1 \times 10^5 \text{ cpm}/\mu\text{g antibody}$
- $1 \times 10^4 \text{ cpm}/\text{ng antibody}$
- Molecular wt. of antibody = 75,000 ng/nmole
- **Specific activity:**
- 25 $1 \times 10^4 \text{ cpm}/\text{ng} \times 75,000 \text{ ng}/\text{nmole} = 7.5 \times 10^8 \text{ cpm}/\text{nmole}$

The antigen-coated plate is blocked (to eliminate non-specific binding, e.g., with mB7.1-CHO) and the background plate (i.e., Untransfected-CHO) for one hour

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at room temperature-with 200 μ l/well of blocking buffer (Blocking Buffer: 1xPBS containing 10% Gelatin, 2% Sodium Azide, 1% BSA and 10% FBS).

5 The plate(s) are then washed, typically ten times by hand with tap water.

The 10 μ g/mL radiolabeled antibody (50 μ ls) is then titrated by two-fold serial dilutions across the plate(s) using a multichannel pipette. Incubate for one hour at room temperature.

10 The plate(s) are again washed about 6-7 times with 200 μ l/well of wash buffer (Wash Buffer:--1xPBS containing 10% Gelatin and 2% Sodium Azide).

The radio activity counts in each well are then determined by running the wells on a gamma counter.

15 The optimal radiolabeled antibody concentration is the concentration in which the difference between the specific counts and background counts is at a maximum.

3. Scatchard Analysis of Competition Assay

20 The 10 μ g/mL radiolabeled solution is diluted to the optimal concentration determined in the **Direct Binding experiment**.

The antigen-coated plate and the background plate are blocked for one hour at room temperature with 200 μ l/well of blocking buffer.

25 The plate(s) are then washed, e.g., about 10 times, by hand with tap water.

30 The "cold" (no radiolabel) antibody is then titrated by two-fold dilutions in a separate U-bottom microtitre plate. The starting concentration of the "cold" antibody should be at least 100 times greater than that of the optimal radiolabeled antibody concentration.

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Example:

- Optimal Radiolabeled Conc.: 0.5 $\mu\text{g/mL}$
- "Cold" Antibody Conc.: 100 $\mu\text{g/mL}$ (Note: 1:2 titration in the first well will adjust the "cold" antibody concentration to 50 $\mu\text{g/mL}$.)

Fifty μl /well of optimal radiolabeled antibody are then added to the wells containing "cold" antibody.

One hundred μl /well of the mixed solution are then transferred to the corresponding wells of the antigen-coated plate, and incubated for one hour at room temperature.

Also, it is desirable also that the following controls be effected:

a) Direct binding of radiolabeled antibody to antigen-coated plate (5 wells),

b) Direct binding of radiolabeled antibody to background plate (5 wells).

After incubation, the plate(s) are washed, e.g., about 6-7 times, with 200 μl /well of wash buffer.

The radio activity counts in each well are then determined by running the wells on a gamma counter.

These calculations are determined by calculating the specific counts in each well tested by subtracting the background counts from the counts bound to the antigen-coated plate.

4. Calculations for Scatchard Analysis

The Molar Concentration of Bound antibody [B] can then be determined as follows:

Example: At 50 $\mu\text{g/mL}$ "cold antibody"

Specific counts bound: 4382 cpm

Counts bound in the presence of 50 $\mu\text{g/mL}$ "cold" ab: 215 cpm

Difference: 4382 cpm - 215 cpm = 4167 cpm

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Specific Activity (radiolabeled ab): 5.54×10^9 cpm/nmole
 $4167 \text{ cpm} \div 5.54 \times 10^9 \text{ cpm/nmole} = 7.52 \times 10^{-7} \text{ nmole}$
 $7.53 \times 10^{-7} \text{ nmole} \div 0.05 \text{ mL (sample vol.)}$
 $= 1.50 \times 10^{-5} \text{ nmole/mL}$
5 $= 1.50 \times 10^{-8} \text{ } \mu\text{mole/mL}$
 [B] $= 1.50 \times 10^{-11} \text{ mole/mL (M)}$

Total Molar Concentration [T] is determined as follows:

$50 \text{ } \mu\text{g/mL} \times 1 \text{ } \mu\text{mole/75,000 } \mu\text{g} = 6.67 \times 10^{-4} \text{ } \mu\text{mole/mL}$
 $= 6.67 \times 10^{-7} \text{ mmole/mL (M)}$
10 [T] $= 66667 \times 10^{-11} \text{ mmole/mL}$
 (M)

Free antibody [F] is determined as follows:

Free Molar Conc. = Total minus Bound
[F] $= (66667 \times 10^{-11}) - (1.50 \times 10^{-11})$
15 $= 66665.5 \times 10^{-11} \text{ mmole/mL (M)}$

Calculate B/F.

Plot B versus B/F on Cricket Graph software.

Activity and Affinity of Anti-Human CD23
Antibodies According to the Invention

20 Four of the five isolated primate anti-human CD23
monoclonal antibodies (B3B11, 2C8, 5E8 and 6G5) were
found to inhibit IgE production in the above-identified
in vitro assay which measures IgE production by IL4-
hydrocortisone induced peripheral blood mononuclear cell
25 (PBMC) cultures. These results are shown in Figure 1.
The fifth primate monoclonal anti-human CD23 antibody
3G12 was inactive in this assay.

Two of the four primate monoclonal anti-human CD23
antibodies (B3B11 and 2C8) found to be active in this in

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vitro assay were found to compete with a commercially available mouse anti-human CD23 antibody MHM6 (CAKO A/S, Glostrup, Denmark Catalog # M763). (Figure 2, top panel.) However, in repeated assays these antibodies were not as potent IgE inhibitors as MHM6 (data not shown). By contrast, the other primate anti-CD23 monoclonal antibodies (5E8 and 6G5) were found to compete with each other and did not compete with MHM6. (Figure 2, middle and bottom panels.) Moreover, the primate anti-human CD23 monoclonal antibody 5E8 was found to be a potent inhibitor of IL-4 induced IgE in the *in vitro* assay. (See Figures 1 and 3.)

Modified Hu-SCID-Mouse Model for Human IgE Synthesis and Measuring the Inhibition of IL-4 Induced IgE Production by Anti-CD23 Antibodies In Vivo

A modified hu-PBMC-SCID mouse model was also developed to detect the effect of the subject antibodies on induced human IgE production *in vivo*. PBMCs obtained from two donors were cultured with IL-4 *in vitro* for two days. PBMCs were pooled and used to reconstitute groups of C.B.-17 SCID mice with and without antibodies. Mice were bled on day 14, 21, 28 and 35 and serum IgG and IgE levels were determined by ELISA. This *in vivo* model was used to assay primate and two different versions of PRIMATIZED® antibodies to CD23 for their ability to inhibit the production of IgE.

A modified SCID mouse model was used because it is known that severe combined immunodeficiency scid/scid (SCID) mice, C.B.-17 (Bosma et al., *Nature*, 301:527 (1983)) reconstituted with human peripheral blood mononuclear cells (hu-PBMC-SCID) can produce significant quantities of human immunoglobulins (Ig) (Moiser et al.,

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Nature, 335:256 (1988); Moiser et al., *J. Clin. Immunol.*, 10:185 (1990); Abedi et al., *J. Immunol.*, 22:823 (1992); and Mazingue et al., *Eur. J. Immunol.*, 21:1763 (1991).) The predominant isotype of human immunoglobulin (Ig) produced in hu-PBMC-SCID mice is IgG. Generally, IgM, IgA and IgE isotypes are found in very low or non-detectable levels except in cases where PBMC is obtained from donors with certain autoimmune or allergic disease conditions. It has also been reported that manipulation of hu-PBMC SCID mouse model with certain cytokines may be provided for the generation of significant levels of non-IgG isotypes, including IgE (Kilchherr et al., *Cellular Immunology*, 151:241 (1993); Spiegelberg et al., *J. Clin. Investigation*, 93:711 (1994); and Carballido et al., *J. Immunol.*, 155:4162 (1995)). The hu-PBMC-SCIDs, has been also used to generate antigen specific Ig provided the donor has been primed for the antigen *in vivo*.

Therefore, the aim of the present inventors was focused on establishing a suitable human IgE producing hu-PBMC-SCID mouse model that could be used to test the efficacy of therapeutic for treatment of IgE related diseases such as allergic disorders, including the subject anti-CD23 antibodies.

Materials and Methods:

The following materials and methods were used in the hu-PBMC-SCID mouse model described below.

SCID mice: C.B-17 scid/scid immunodeficient mice were obtained from Taconic (C.B. -17/IcrTac-scidfDF) and maintained in IDEC Pharmaceuticals' animal facility. Mice were housed in sterilized microbarrier units with sterilized bedding. Animal studies were performed in

accordance with the "Guide for the Care and Use of Laboratory Animals" specified by the Committee on Care of Laboratory Animal Resources Commission on Life Science-National Research Council (Guide for the Care and Use of Laboratory Animals, DHHS Publ. No. (NIH) 86-23, Bethesda, MD, NIH, 1985).

Human PBMC: PBMCs were isolated from buffy coats obtained from a blood bank by centrifugation through Ficoll-Hypaque (Histopaque-1077) as recommended by the manufacturer (Sigma Diagnostics Catalog # 1077-1). Lymphocyte preparation at the interface of the gradient were harvested and washed three times in Hanks Balanced Salt Solution (HBSS) (Bio-Whittaker Catalog # 10-527F). For each experiment PBMCs were obtained from two separate donors and cultured separately *in vitro*. PBMCs were resuspended at $1-3 \times 10^6$ cells/ml concentration in HB-Basal medium plus 1% HB101 lyophilized supplement (Irvine Scientific Catalog # T000 & T151) containing 5% FCS plus 1000 IU/ml of IL-4 (Genzyme, Inc. Catalog # 2181-01) and incubated for 48 hours at 37°C with 5% CO₂. After incubation, the cells from different buffy coats were harvested, pooled and used to reconstitute SCID mice.

In Vivo Assay Conditions

Groups of mice (four to five per group) were injected with fifty-sixty $\times 10^6$ lymphocytes in 200-300 μ l volume of HBSS intraperitoneally (i.p.) on day zero. For the groups that received anti-CD23 antibody, on day zero, PBMCs were mixed with anti-CD23 antibody (200 to 400 μ g/mouse) before i.p., injection and the second injection was given on day seven. All mice received 5000 IU per mouse of IL-4 i.p., between day zero to day five. A group which was not injected with antibody

served as the control group. Mice were bled from a retro-orbital vein and the serum was analyzed for IgG and IgE on days fourteen, twenty-one, twenty-eight and thirty-five by ELISA.

- 5 Figure 8 shows that the primate anti-human CD23 monoclonal antibody 5E8 is effective in inhibiting IL-4 induced IgE production *in vivo* in the SCID mouse model.

**Cloning and Expression of PRIMATIZED® anti-human
CD23 Monoclonal Antibodies**

- 10 In order to clone primate immunoglobulin variable domains, Poly A+ RNA was separately isolated from approximately 2×10^6 cells from the primate heterohybridomas secreting the anti-human CD23 monoclonal antibodies 6G5 and 5E8 by using the
15 Micro-FastTrack mRNA isolation Kit (Invitrogen Catalog # K1520-02) according to methods set forth by the manufacturer.

- The first strand of cDNA was synthesized from the poly A+ RNA by using the cDNA Cycle Kit (Invitrogen
20 Catalog # L1310-01) according to conventional methods.

- The light and heavy chain variable regions of 6G5 and 5E8 were then isolated by PCR from cDNA using PCR primers that were selected based upon different consensus families of human immunoglobulins. 5' primers
25 were selected which corresponded to the beginning of the leader sequences of the light and heavy variable region and 3' primers were selected which corresponded to the J region (The specific primers used to PCR amplify the lambda light chain variable domain of 6G5, the kappa
30 light chain variable domains of 5E8, and the heavy chain variable domains of 6G5 and 5E8 are set forth in Tables 1-3). PCR was performed according to standard methods (30 cycles with 1 minute at 94°C, 1.5 minutes at 54°C and

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2 minutes at 72°C in a Hot start 100 tube (Gibco BRL Catalog # 10332-013). PCR was set up in 50µl reactions containing 5µl out of 80µl cDNA (from 2x10⁶ cells) as a template, 2µl of 5 nM dNTP, 1µl of Taq polymerase, 5µl of Taq polymerase buffer, 2µl of the 5' primer (25 pmoles/µl), 2µl of the 3' primer (25 pmoles/µl), and 36µl of water. (Taq polymerase and buffer were obtained from Stratagene Catalog # 600131, dNTP from Boehringer Mannheim Catalog #1581295.)

10 A) Construction of the plasmids N5LG1 + 6G5 and N5LG4P + 6G5

1) Cloning the light chain variable domain of primate monoclonal anti-human CD23 antibody 6G5 by PCR

15 The first PCR amplification of the light chain variable region from the cDNA of primate monoclonal antibody 6G5 showed bands which were consistent in situ with the lambda light chain variable region. These bands appeared in all reactions using the three different early leader sequence primers. (See Tables 1-3.) However, the PCR product obtained using primer 745 (Family 2) was considered more specific because of the relatively greater intensity of the PCR product band.

20 This PCR product was isolated using a Qiaquick Gel Extraction Kit (Qiagen Catalog # 28704). The purified PCR fragment was digested with Bgl II and Avr II restriction endonucleases, and ligated into the mammalian expression vector N5LG1 which was digested with the same restriction endonucleases. Twenty microliters of the ligation mixture containing the purified PCR product from one fifty microliter PCR reaction, 100 mg N5LG1 vector, two microliters of 10x ligation buffer (NEB Catalog # 202S) and two microliters of T4

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ligase (NEB Catalog # 202S), were then incubated at 14°C overnight.

The mammalian expression vector N5LG1 contains genetic sequences (e.g., regulatory sequences, coding sequences) which provide for the expression of four separate proteins in a mammalian cell. They are:

(i) a partial immunoglobulin light chain with the human lambda light chain constant region and unique restriction endonuclease sites for inserting light chain variable domains;

(ii) a partial immunoglobulin heavy chain with the and human gamma 1 chain constant region coding sequences and unique restriction endonuclease sites for inserting heavy chain variable domains;

(iii) a neomycin phosphotransferase gene used to select for cells that have incorporated the plasmid and are resistant to the antibiotic Geneticin (Gibco BRL Catalog # 10131-1209); and

(iv) a murine dihydrofolate reductase gene (DHFR) which provides for the selection and genomic amplification when cells are cultured in the presence of methotrexate (MTX, Sigma Catalog # A-6770) (Reff et al., *Blood*, 83:433-445 (1994)).

After ligation, the mixture was digested using Pme I restriction endonuclease, which digests the parent N5LG1 plasmid, but not the N5LG1 plasmid which has been ligated to the light variable domain of 6G5. After digestion, the mixture was transformed into Epicurian coli® XL1-Blue competent cells (Stratagene Catalog # 200249) as follows.

One hundred microliters of competent cells were mixed with 10 µl of the above ligation mixture, set on ice for 30 minutes, then heated at 45°C for 30 seconds. This mixture was placed on ice for 2 minutes, and 900 µl

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of SOC, prewarmed to room temperature, was then added. (SOC is LB broth Gibco BRL Catalog # 10855-013, plus 0.02 M $MgCl_2$, 0.02 M $MgSO_4$ and 0.02 M D-glucose.) After incubation at 37°C for an hour, the mixture was

5 centrifuged at 4000 g for a minute, and 800 μ l of supernatant discarded. The rest of the mixture was plated onto a LB agar (Gibco BRL Catalog # 12945-044) dish containing 50 μ g/ml ampicillin (Amp, Gibco BRL Catalog # 13075-015). Plasmid DNA was isolated from

10 individual colonies of *E. coli* that grew on the Amp plate by using the Wizard® Miniprep DNA purification system (Promega Catalog # A7510).

The isolated plasmid DNA was then characterized by digestion with Bgl II and Avr II followed by agarose gel

15 electrophoresis. An ethidium stained DNA band of 400 bp was indicative of a potential successful cloning of a light chain variable domain.

To confirm this was an immunoglobulin light chain variable domain, sequencing was done using the Sequenase

20 7-Deaza -dGTP DNA Sequencing Kit (USB catalog # 70990) with sequencing primers 607 and GE 108. (See Sequencing Primers in Table 4.)

A second independent PCR amplification of the light chain from cDNA of primate monoclonal antibody 6G5 was

25 effected using a 5' primer early leader sequence of lambda light chain family 2 (primer 745) and the 3' J region primer 926. (See Primers for PCR of the lambda light chain variable domain of 6G5 in Tables 1-3.) The isolated PCR product (see technique above) was cloned

30 into TA vector by using the Original TA Cloning(Kit (Invitrogen Catalog # K2000-01). The isolated miniprep DNA (see technique above) was examined under agarose gel electrophoresis after digestion with EcoR I restriction endonuclease. The resultant PCR product comprised in

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the TA vector was then sequenced (as described previously) using Sp6 and M13(-40) forward primers (See Sequencing primers in Table 4). The resultant light chain sequence was identical to that of light chain from the first PCR. This entire sequence of the light chain variable domain of primate monoclonal anti-human CD23 antibody 6G5 is presented below.

SEE ID NO: 11

Light chain variable region of primate monoclonal antibody anti-human CD23 6G5 Leader

10 Met Ala Trp Thr Leu Leu Leu Val Thr Leu Leu Thr Gln Gly Thr
ATG GCC TGG ACT CTG CTC CTC GTC ACC CTC CTC ACT CAG GGC ACA
-1
Gly Ser Trp Ala
GGA TCC TGG GCT

15 Mature Protein (Numbering is Kabat)

Framework 1

1 9 11
Gln Ser Ala Pro Thr Gln Pro Pro Ser Val Ser Gly Ser Pro Gly
CAG TCT GCC CCG ACT CAG CCT CCC TCT GTG TCT GGG TCT CCT GGA
20 20 23
Gln Ser Val Thr Ile Ser Cys
CAG TCG GTC ACC ATC TCC TGC

CDR 1

24 27 27A 27B 27C 28 34
25 Thr Gly Thr Ser Asp Asp Val Gly Gly Tyr Asn Tyr Val Ser
ACT GGA ACC AGC GAT GAC GTT GGT GGT TAT AAC TAT GTC TCC

Framework 2

35 40 49
30 Trp Tyr Gln His His Pro Gly Lys Ala Pro Lys Leu Met Ile Tyr
TGG TAC CAA CAC CAC CCA GGC AAA GCC CCC AAA CTC ATG ATT TAT

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CDR2

50 56
Asp Val Ala Lys Arg Ala Ser
GAT GTC GCT AAG CGG GCC TCA

5

Framework 3

57 60 70
Gly Val Ser Asp Arg Phe Ser Gly Ser Lys Ser Gly Asn Thr Ala
GGG GTC TCT GAT CGC TTC TCT GGC TCC AAG TCT GGC AAC ACG GCC

10

80
Ser Leu Thr Ile Ser Gly Leu Gln Ala Glu Asp Glu Ala Asp Tyr
TCC CTG ACC ATC TCT GGG CTC CAG GCT GAG GAC GAG GCT GAT TAT

88
Tyr Cys
TAC TGT

15

CDR 3

89 90 95 95A 96 97
Cys Ser Tyr Thr Thr Ser Ser Thr Leu Leu
TGT TCA TAT ACA ACC AGT AGC ACT TTG TTA

Framework 4

20

98 100 106 106A 107
Phe Gly Arg Gly Thr Arg Leu Thr Val Leu Gly
TTC GGA AGA GGG ACC CGG TTG ACC GTC CTA GGT

2) Cloning the heavy chain variable domain of primate monoclonal anti-human CD23 antibody 6G5 by PCR

25

The first PCR amplification of the heavy chain variable domain from cDNA of primate monoclonal antibody 6G5 was performed by using the set of early leader sequence primers described supra and the 3' J region primer GE244. These primers are in Tables 1-3 infra.

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1 This reaction resulted in a 350 base PCR product. This
2 350 base product (purified as described supra), was di-
3 gested with Nhe I and Sal I, and ligated into N5LG1 and
4 digested with the same endonucleases in the first PCR
5 amplification. The resultant ligation mixture was
6 transformed into host cells using the same techniques
7 for cloning the light chain. Plasmid N5LG1 containing
8 the 350 base PCR product was then isolated and sequenced
9 (using sequencing primers 266 and 268). (These
10 Sequencing primers are set forth in Table 4.)

11 Sequencing revealed that the PCR product contained
12 only part of the heavy variable domain and comprised a
13 deletion in its amino terminus (Sequence began at frame-
14 work 2, codon 36).

15 A second independent PCR reaction was conducted to
16 amplify and isolate the heavy chain variable domain of
17 primate monoclonal antibody 6G5 using a 5' early leader
18 sequence primer for family 1 (MB1503) and a 3' J' region
19 primer GE244. (These primers are also contained in
20 Tables 1-3.) The resultant PCR product was then cloned
21 into the N5LG1 using the same techniques described
22 supra. Its sequence was found to be identical to the
23 first PCR product.

24 Therefore, in order to clone the whole heavy
25 variable domain of 6G5 including the missing 5' terminus
26 a new longer 3' primer (MB1533) which included the CDR3
27 and framework 4 regions of the 6G5 heavy variable chain
28 was then used in a third independent PCR reaction with
29 the family 1 5' primer (MB1503). (These primers are
30 also contained in Tables 1-3.)

After PCR, a larger 420 base PCR product was
observed on the agarose gel. This PCR product was
isolated as described previously, and cloned into a TA
vector. The resultant PCR product contained in the TA

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vector was then sequenced. Sequencing revealed that this DNA contained the whole heavy variable domain and that the 3' part was identical to that of previously cloned partial heavy chain variable domain from the first two PCR reactions.

A fourth independent PCR was performed using the same primers as the third PCR amplification. This resulted in a PCR product which was isolated and cloned into the TA vector as described previously. The sequence of the fourth independent PCR product was found to be identical to that obtained in the third PCR amplification. This sequence, which comprises the heavy chain variable domain of primate monoclonal anti-human CD23 antibody 6G5, is presented below

Heavy chain variable region of primate monoclonal antibody anti-human CD23 6G5

Leader

Met Lys His Leu Trp Phe Phe Leu Leu Leu Val Ala Ala Pro Arg
ATG AAA CAC CTG TGG TTC TTC CTC CTC CTG GTG GCA GCT CCC AGA
-1
Trp Val Leu Ser
TGG GTC CTG TCC

Mature Protein (Numbering is Kabat)

Framework 1

1 10
Gln Leu Gln Leu Gln Glu Ser Gly Pro Gly Val Val Lys Pro Ser
CAG CTG CAG CTG CAG GAG TCG GGC CCA GGA GTG GTG AAG CCT TCG
20 30
Glu Thr Leu Ser Leu Thr Cys Ala Val Ser Gly Gly Ser Val Ser
GAG ACC CTG TCC CTC ACC TGC GCT GTC TCT GGT GGC TCT GTC AGC

CDR 1

31 35 35a
Ser Ser Asn Trp Trp Thr
AGT AGT AAC TGG TGG ACC

5

Framework 2

36 40 49
Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile Gly
TGG ATC CGC CAG CCC CCA GGG AAG GGA CTG GAG TGG ATT GGA

CDR2

10

50 52 52A 53 60
Arg Ile Ser Gly Ser Gly Gly Ala Thr Asn Tyr Asn Pro Ser Leu
CGT ATC TCT GGT AGT GGT GGG GCC ACC AAC TAC AAC CCG TCC CTC
65

Lys Ser

15

AAG AGT

Framework 3

66 70 80
Arg Val Ile Ile Ser Gln Asp Thr Ser Lys Asn Gln Phe Ser Leu
CGA GTC ATC ATT TCA CAA GAC ACG TCC AAG AAC CAG TTC TCC CTG

20

82 82a 82b 82c 83 90
Asn Leu Asn Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys
AAC CTG AAC TCT GTG ACC GCC GCG GAC ACG GCC GTG TAT TAC TGT

94

Ala Arg

25

GCC AGA

CDR 3

95 100 100a 100b 100c 100d 101 102
Asp Trp Ala Gln Ile Ala Gly Thr Thr Leu Gly Phe
GAT TGG GCC CAA ATA GCT GGA ACA ACG CTA GGC TTC

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365020-TH06060

103							110			113
Trp	Gly	Gln	Gly	Val	Leu	Val	Thr	Val	Ser	Ser
TGG	GGC	CAG	GGA	GTC	CTG	GTC	ACC	GTC	TCC	TCA

In order to insert the cloned heavy chain variable domains of 6G5 into a mammalian expression vector, the heavy chain variable domain in the TA vector (obtained in the 3rd independent PCR) was digested with Nhe I and Sal I and cloned into the N5LG1 vector which was digested with the same restriction enzymes and which vector already contains the light chain variable domain. The resultant mammalian expression vector was named N5LG1 + 6G5.

15 To construct the N5LG4P + 6G5 vector, both the light and heavy chain variable domains were isolated from N5LG1 + 6G5 by digestion of Bgl II and Avr II, and Nhe I and Sal I respectively. The mammalian expression vector N5LG4P vector is identical to the N5LG1 vector
20 described above, except the human gamma 1 was replaced with a human gamma 4 constant region containing a mutation of a serine to a proline in the hinge region to increase stability of the immunoglobulin and improve pharmacokinetics *in vivo* ("P" mutation). The light
25 chain variable domain was cloned in the plasmid first and the heavy chain variable domain was cloned into the vector containing the light chain variable domain using techniques previously described. This mammalian expression vector was named N5LG4P + 6G5.

B. Construction of the plasmids N5KG4P + 5E8, N5KG1 + 5E8, N5KG4P + 5E8N-, and N5KG1 + 5E8N-

1. Cloning the light chain variable domain of primate monoclonal anti-human CD23 antibody 5E8 by PCR

5 The first PCR reaction of the light chain variable domain from 5E8 cDNA was carried out using a set of kappa early leader sequence primers and the 3' J region primer GE204. (See primers for PCR of the kappa light chain variable domain of 5E8 in Tables 1-3). A 420 base
10 PCR product was obtained. The isolated 420 base PCR product was digested with Bgl II and BsiW I restriction endonucleases, cloned into the mammalian expression vector N5KG4P and sequenced using GE108 and 377 primers (which are contained in Table 4). The mammalian
15 expression vector N5KG4P is identical to the vector N5LG4P except it contains the human kappa light chain constant region in place of the human lambda light chain constant region. Sequencing of this 420 polynucleotide DNA revealed that it contains the entire
20 kappa light chain variable domain.

 A second independent PCR of the light chain variable region was performed using the 5' family 1 primer GE201 and the 3' primer GE204. (See primers for PCR of the kappa light chain variable domain of 5E8 in
25 Tables 1-3). The isolated PCR product was cloned into the TA vector (using methods previously described) and sequenced using Sp6 and T7 promoter primers. Sequencing revealed that this PCR product was identical to that obtained from the first PCR. The entire sequence of the
30 light chain variable domain of primate monoclonal anti-human CD23 antibody 5E8 is presented below.

5E8 ID NO:3

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B

Light chain variable region of primate monoclonal
antibody anti-human CD23 5E8

Leader

Met Asp Met Arg Val Pro Ala Gln Leu Leu Gly Leu Leu Leu Leu
5 ATG GAC ATG AGG GTC CCC GCT CAG CTC CTG GGG CTC CTT CTG CTC

-1

Trp Leu Pro Gly Ala Arg Cys
TGG CTC CCA GGT GCC AGA TGT

Mature Protein (Numbering is Kabat)

10

Framework 1

1 10
Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val
GAC ATC CAG ATG ACC CAG TCT CCA TCT TCC CTG TCT GCA TCT GTA

20

23

15

Gly Asp Arg Val Thr Ile Thr Cys
GGG GAC AGA GTC ACC ATC ACT TGC

CDR 1

20

24 30 34
Arg Ala Ser Gln Asp Ile Arg Tyr Tyr Leu Asn
AGG GCA AGT CAG GAC ATT AGG TAT TAT TTA AAT

Framework 2

25

35 40 49
Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr
TGG TAT CAG CAG AAA CCA GGA AAA GCT CCT AAG CTC CTG ATC TAT

CDR2

50 56
Val Ala Ser Ser Leu Gln Ser
GTT GCA TCC AGT TTG CAA AGT

Framework 3

30

57 60 70
Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Glu Phe
GGG GTC CCA TCA AGG TTC AGC GGC AGT GGA TCT GGG ACA GAG TTC

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80

Thr Leu Thr Val Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr
ACT CTC ACC GTC AGC AGC CTG CAG CCT GAA GAT TTT GCG ACT TAT

88

5 Tyr Cys
TAC TGT

CDR 3

89 90

97

10 Leu Gln Val Tyr Ser Thr Pro Arg Thr
CTA CAG GTT TAT AGT ACC CCT CGG ACG

Framework 4

98

100

107

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
TTC GGC CAA GGG ACC AAG GTG GAA ATC AAA

15 2) Cloning the heavy chain variable domain of primate
monoclonal anti-human CD23 antibody 5E8 by PCR

20 The first PCR of the heavy chain variable domain of
5E8 was performed using a set of 5' early leader heavy
chain sequence primers and the 3' primer GE210. (See
primers for PCR of the heavy chain variable domain of
6G5 and 5E8 in Table 1). A 420 base PCR product
appeared in the family 3 primer reaction. The PCR
product was purified and then digested with Nhe I and
Sal I and cloned into the mammalian expression vector
25 N5KG4P vector (as described previously). The PCR
product was sequenced using the 268 and 928 primers.
(See sequencing primers in Table 4.)

30 A second independent PCR of the heavy chain
variable domain of 5E8 was performed using the family 3
5' primer GE207 and the 3' primer GE210. (See primers
for PCR of the heavy chain variable domain of 6G5 and

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5E8 in Tables 1-3). The isolated PCR product was cloned into a TA vector using the same techniques previously described and sequenced by using Sp6 and T7 primers. Sequencing revealed that the TAC at codon 91 had been changed into TGC.

In order to determine the appropriate codon at 91, a third independent PCR was performed using the same primers as the second PCR (see above). The PCR product was again cloned into a TA vector and sequenced using Sp6 and T7 primers. The sequence was found to be identical to the heavy chain variable sequence obtained in the first PCR. Therefore, the TGC at position 91 in the second independent PCR product is apparently the result of an error introduced during PCR. This entire sequence of the heavy chain variable domain of primate monoclonal anti-human CD23 antibody 5E8 is presented below.

Heavy chain variable region of primate monoclonal antibody anti-human CD23 5E8

Leader

Met Glu Phe Gly Leu Ser Trp Val Phe Leu Val Pro Leu Leu Lys
ATG GAG TTT GGG CTG AGC TGG GTT TTC CTT GTT CCT CTT TTG AAA

-1

Gly Val Gln Cys

GGT GTC CAG TGT

Mature Protein (Numbering is Kabat)

Framework 1

1

10

Glu Val Gln Leu Val Glu Ser Gly Gly Gly Leu Ala Lys Pro Gly
GAG GTG CAG CTG GTG GAG TCT GGG GGC GGC TTG GCA AAG CCT GGG

20

30

Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Arg Phe Thr

GGG TCC CTG AGA CTC TCC TGC GCA GCC TCC GGG TTC AGG TTC ACC

CDR 1

31 35 35a 35b
Phe Asn Asn Tyr Tyr Met Asp
5 TTC AAT AAC TAC TAC ATG GAC

Framework 2

36 40 49
Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Val Ser
10 TGG GTC CGC CAG GCT CCA GGG CAG GGG CTG GAG TGG GTC TCA

CDR2

50 52 52A 53 60
Arg Ile Ser Ser Ser Gly Asp Pro Thr Trp Tyr Ala Asp Ser Val
CGT ATT AGT AGT AGT GGT GAT CCC ACA TGG TAC GCA GAC TCC GTG
65
15 Lys Gly
AAG GGC

Framework 3

66 70 80
Arg Phe Thr Ile Ser Arg Glu Asn Ala Asn Asn Thr Leu Phe Leu
20 AGA TTC ACC ATC TCC AGA GAG AAC GCC AAC AAC ACA CTG TTT CTT
82 82a 82b 82c 83 90
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
CAA ATG AAC AGC CTG AGA GCT GAG GAC ACG GCT GTC TAT TAC TGT
94
25 Ala Ser
GCG AGC

CDR 3

95 100 101
Leu Thr Thr Gly Ser Asp Ser
30 TTG ACT ACA GGG TCT GAC TCC

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103							110		113	
Trp	Gly	Gln	Gly	Val	Leu	Val	Thr	Val	Ser	Ser
TGG	GGC	CAG	GGA	GTC	CTG	GTC	ACC	GTC	TCC	TCA

The heavy variable domain in N5KG4P was digested with Nhe I and Sal I, purified, and cloned into N5KG4P which contains the light chain variable domain of 5E8. This plasmid was then digested with the restriction endonucleases as previously described. This resulted in a vector containing both the light and heavy variable domain of 5E8. This vector was named N5KG4P + 5E8. The heavy and light variable domains of N5KG4P + 5E8 were then both inserted into the mammalian expression vector N5KG1 to create the N5KG1+ 5E8 vector.

Based upon the sequence of 5E8 heavy variable domain, there is a potential glycosylation site of the immunoglobulin at asparagine codon 75. This potential glycosylation site corresponds to a conserved asparagine-linked glycosylation motif having the following tripeptide sequence: (Asp) - (Any amino acid except proline) - (Serine or threonine). Therefore, a glycosylation mutant of 5E8, which would be unable to be glycosylated at this position because of modification of this glycosylation motif, was generated by replacing the asparagine codon 75 with a lysine (which is found in many human immunoglobulins at this position). Site

specific mutagenesis was effected by the following methods.

5 A first PCR was done using N5KG4P + 5E8 as a template and a 3' primer (corresponding to codon 71 to 79) and which contains a mutation at codon 75 (AAC changed to AAG, Primer MB1654, and a 5' primer at the beginning of the leader sequence (Primer MB1650). (See PCR Primers Used for the Generation of a Glycosylation Mutant of the Heavy Chain Variable Region 5E8 set forth in Table 5).

10 A second PCR was performed on the same template by using a 5' primer (corresponding to codon 71 to 79) containing the same mutation (Primer MB1653) and a 3' primer from the end of framework 4 (Primer MB1651) (See PCR Primers Used for the Generation of a Glycosylation Mutant of the Heavy Chain Variable Region of 5E8 in Table 5.)

15 These two PCR products were isolated and mixed in equal molar ratios. A third independent PCR was then carried out by using the mixture of the first and second PCR products as a template with a 5' primer used in the first PCR (MB1650) and a 3' primer used in the second PCR (MB 1651). (See PCR Primers Used for the Generation of a Glycosylation Mutant of the Heavy Chain Variable Region in Table 5.) The PCR product obtained in third PCR was found to contain the heavy variable domain coding region of 5E8 wherein the asparagine 75 had been changed to lysine.

20 The third PCR product was purified, digested with restriction endonucleases Sal I and Nhe I, and cloned into the N5KG4P containing the light variable domain only. The PCR product was sequenced to confirm that it comprised the mutant heavy variable domain. This mammalian expression plasmid was named N5KG4P + 5E8N-.

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To construct of N5KG1 + 5E8N-, both light and heavy variable domains from N5KG4P - 5E8N- were digested with Bgl II and BsiW I, and Nhe I and Sal I respectively. The light variable domain was cloned first and then the heavy variable domain was inserted into the mammalian expression plasmid N5KG1, to create the plasmid N5KG1 + 5E8N-.

Table 1

Primers for PCR of the kappa light chain variable domain of 5E8

NAME	Light chain Vk -early leader 5' (Bgl II)													FAMILY			
						-22	-21	-20	-19	-18		-17	-16	-15	-14		
GE201	5'	AT	CAC	<u>AGA TCT</u>	CTC	ACC		ATG	GAC	ATG	AGG	GTC	CCC	GCT	CAG	3'	1
GE200	5'	AT	CAC	<u>AGA TCT</u>	CTC	ACC				ATG	AGG	CTC	CCT	GCT	CAG	3'	2
GE202	5'	AT	CAC	<u>AGA TCT</u>	CTC	ACC				ATG	GAA	(A/G)CC	CCA	GC(T/G)	CAG		3'
	3																
GE203	5'	AT	CAC	<u>AGA TCT</u>	CTC	ACC				ATG	GTG	TTG	CAG	ACC	CAG	GTC	
	3'4																

Light chain Vk-3' primer (BsiW I)

				113	112	111	110	109	108	107	106	105	104	103
GE204	5'	GG	TGC	AGC	CAC	CGT	AGC	TTT	GAT	(C/T)TC	CA(G/C)			
	CTT	3'												

Table 2

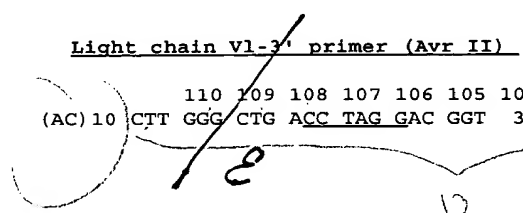
Primers for PCR of the lambda light chain variable domain of 6G5

30

NAME	<u>Light chain VL-early leader 5' (Bgl II)</u>														FAMILY
					-20	-19		-18		-17		-16		-15	
744	5'	AT	CAC	<u>AGA TCT</u>	CTC	ACC	ATG	(G/A)CC	TG(G/C)	TCC	CCT	CT	3'		1
745	5'	AT	CAC	<u>AGA TCT</u>	CTC	ACC	ATG	GCC	TGG	(A/G)CT	C(T/C)G	CT	3'		2
910	5'	AT	CAC	<u>AGA TCT</u>	CTC	ACC	ATG	GC(A/C)	TGG	A(T/C)C	CCT	CTC	3'		3

Light chain VL-3' primer (Avr II)

				110	109	108	107	106	105	104
926	5'	(AC)10	CTT	GGG	CTG	ACC	TAG	GAC	GGT	3'



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Table 3

Primers for PCR of the heavy chain
variable domains from 6G5 and 5E8

NAME	Heavy chain-early leaders 5' (Sal I)	Family
5		
	-20 -19 -18 -17 -16 -15	
MB1503	5' GCG ACT AAG <u>TCG ACC</u> ATG GAC TGG ACC TGG 3'	1 ID NO 114
MB1502	5' GCG ACT AAG <u>TCG ACC</u> ATG AAA CAC CTG TGG 3'	2,4 ID NO 115
GE207	5' GCG ACT AAG <u>TCG ACC</u> ATG GAG TTT GGG CTG AGC 3'	3 NO 116
GE208	5' GCG ACT AAG <u>TCG ACC</u> ATG GGG TCA ACC GCC ATC 3'	5 NO 117
10	GE209 5' GCG ACT AAG <u>TCG ACC</u> ATG TCT GTC TCC TTC CTC 3'	6 NO 118
	<u>Heavy chain-3' primer (Nhe I)</u>	
	120 119 118 117 116 115 114 113 112 111 110	
GE244	5' GC CAG GGG GAA GAC CGA TGG GCC CTT GGT <u>GCT AGC</u> TGA GGA GAC GG 3'	NO 119
GE210	5' GA TGG GCC CTT GGT <u>GCT AGC</u> TGA GGA GAC GG 3'	NO 120
15	MB1533 5' GGT <u>GCT AGC</u> TGA GGA GAC GGT	
	109 108 107 106 105 104 103 101 100 99	
	GAC CAG GAC TCC CTG GCC CCA GAA GCC TAG 3'	ID NO 121

Table 4

Sequencing Primers

20	Sp6 primer (SEQ ID NO: 22) 5' AT TTA GGT GAC ACT ATA	3' NO 22
	M13 (-40) Forward	
	Primer (SEQ ID NO: 23) 5' GTT TTC CCA GTC ACG A	3' NO 23
	(SEQ ID NO: 24)	
	T7 Promoter Primer 5' AT ATA CGA CTC ACT ATA GGG	3' NO 24
25	GE 108 Primer (SEQ ID NO: 25) 5' CCG TCA GAT CGC CTG GAG ACG CCA 3'	NO 25
	377 Primer (SEQ ID NO: 26) 5' GCA GTT CCA GAT TTC AAC TG	3' NO 26
	607 PRIMER (SEQ ID NO: 27) 5' CCA GGC CAC TGT CAC GGC TTC	3' NO 27
	266 PRIMER (SEQ ID NO: 28) 5' CAG AGC TGG GTA CGT CCT CA	3' NO 28
	268 PRIMER (SEQ ID NO: 29) 5' GCC CCC AGA GGT GCT CTT GG	3' NO 29

876 PRIMER (SEQ ID NO: 30) 5' ACA CAG ACC CGT CGA CAT GG 3' NO: 30
928 PRIMER (SEQ ID NO: 31) 5' GCT CTC GGA GGT GCT CCT GG 3' NO: 31

Table 5

5 PCR Primers Used for the Generation of a Glycosylation
Mutant of the Heavy Chain Variable Region of 5E8

Sal I -20 -19 -18 -17 -16
(SEQ ID NO: 32)
10 MB 1650_A 5' ACA GAC CCG TCG ACC ATG GAG TTT GGG CTG 3' (SEQ ID NO: 32)
Nhe I
(SEQ ID NO: 33) 118 117 116 115 114 113 112 111 110
MB 1651_A 5' CCC CTT GGT GCT AGC TGA GGA GAC GGT 3' (SEQ ID NO: 33)
(SEQ ID NO: 34) 71 72 73 74 75 76 77 78 79
15 MB 1653_A 5' AGA GAG AAC GCC AAG AAC ACA CTG TTT 3' (SEQ ID NO: 34)
(SEQ ID NO: 35) 79 78 77 76 75 74 73 72 71
MB 1654_A 5' AAA CAG TGT GTT CTT GGC GTT CTC TCT 3' (SEQ ID NO: 35)

C) Construction of gamma-3 vectors

Numerous methods exist for conversion of murine
antibodies to chimeras in which the heavy and light
20 chain constant regions are substituted with human
versions or in which all but the CDRs (complementary
determining regions) are substituted with their human
equivalents. (See Biochem. J. 281:317, 1992; Proc. Nat.
Acad. Sci. USA 86:10029, 1989; Methods Enzymol. 178:515,
25 1989; Cancer Res. 51:181, 1991; Biotechniques 7:360,
1989; J. Immunol. 143:3589, 1989; Int. J. Cancer 44:424,
1989; Proc. Nat. Acad. Sci. USA 86:3833, 1989).

Further, the previous sections specifically
demonstrate that construction of vectors for expression
30 of PRIMATIZED® gamma-1 and gamma-4 antibodies may be
accomplished by amplifying the DNA encoding the light

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and heavy chain variable regions using PCR, and cloning these regions into an expression vector or vectors encoding the appropriate human constant region domains. Thus, it should be clear that similar constructs may be made which encode other human constant region domains, so long as appropriate mammalian expression vectors are available which also contain cloning sites which allow insertion of the amplified variable region DNA from the primate antibodies.

It should also be apparent that the PCR primers used to amplify the variable region DNA may be designed to accommodate various restriction cloning sites depending on the vector used.

Vectors encoding the constant region domain from human IgG3 are available in the art, and may be used for constructing the PRIMATIZED® antibodies of the present invention. For instance, Co et al. have used separate vectors expressing human light and heavy chain constant regions for constructing chimeric and humanized antibodies, and provide heavy chain vectors for both human gamma-1 and gamma-3. For cloning the variable domain regions of the antibody of interest into these vectors, an *Xba*I restriction site is conveniently located just before the constant regions in the J_κ4 and the J_H3 intronic segments for the light and heavy chains, respectively (Co et al., 1996, Cancer Res. 56: 1118-1125; Co et al., 1992, J. Immunol. 148: 1149-1154).

Thus, by designing PCR primers which incorporate *Xba*I restriction sites, or sites which create the same single stranded overhangs as the *Xba*I restriction enzyme (i.e. *Nhe*I, *Spe*I), the variable regions from a primate antibody may be associated with the human gamma-3 constant region domain using the vectors described in

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Co et al. Alternatively, linkers or adaptors may be used to facilitate cloning.

It is also possible to construct a cDNA version of the human gamma-3 constant region using DNA synthesis techniques and insert it into the N5KG1 vectors described herein in place of the human gamma-1 constant region between the *NheI* and *BamHI* sites. Depending on which restriction sites are most convenient for cloning the particular variable region of interest, there are also other gamma-3 vectors which have been described in the art, and any of these may be used to construct and express the gamma-3 antibodies of the present invention. (See, for example, Parren et al., 1992, J. Immunol. 148(3): 695-701; Steplewski et al., 1988, Proc. Natl. Acad. Sci. USA 85: 4852-4856; and U.S. Patent No. 4,975,369, herein incorporated by referenced).

D) Variations of PRIMATIZED® Antibodies

The present invention also encompasses PRIMATIZED® antibodies containing mutations, substitutions or deletions of the constant region. Such mutations may be constructed using site-directed mutagenesis techniques as described above for the glycosylation mutant, which are well-known techniques for those of skill in the art.

The mutated antibodies of the present invention are designed with the purpose of creating a desired change in the level of therapeutic efficiency, i.e. in FcR binding. However, it should be understood that any mutation in the constant region of the antibodies should not alter the basic effector functions as mediated by the gamma-1 or gamma-3 constant region domains. It may also be possible to alter other subclasses of gamma antibodies by mutagenesis such that they have similar effector functions as an antibody containing a gamma-1

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or a gamma-3 domain. Such antibodies are also encompassed by the present invention.

For example, the regions of the IgG constant region involved in FcR binding and interaction with the C1q complement component have been characterized, and
5 mutations have been identified which either increase or decrease binding affinity. As disclosed in U.S Patent 5,648,260, herein incorporated by reference, changing Leu 235 to Glu in the human IgG3 constant region
10 destroys the interaction of the mutant for the human Fc gamma R1 receptor. Furthermore, mutations on adjacent or close sites in the hinge link region (i.e. replacing residues 234, 236 or 237 by Ala) indicate that alterations in residues 234, 235, 236 and 237 at least
15 affect affinity for the Fc gamma R1 receptor.

Similarly, in U.S. Patent 5,348,876, also incorporated by reference, it was shown that by decreasing the length of the hinge region of the human IgG3 constant region and altering the amino acid
20 sequence of this region, C1q binding and thus complement mediated cytolysis were further improved. In fact, the variants bound more C1q than wild type IgG3 and much more than IgG1.

Thus, it is clear from the art that mutations may
25 be used to create therapeutic antibodies with decreased affinities for human Fc receptors, perhaps for the treatment of more mild medical conditions. In addition, mutations may be identified which increase effector function activity, thereby leading to stronger versions
30 for therapeutic use. It should be recognized that manipulations of gene sequence to create alterations in drug efficiency are envisioned, and well within the spirit and scope of the present invention.

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E) Expression of PRIMATIZED® Antibodies
in Chinese Hamster Ovary Cells

In order to isolate the PRIMATIZED® antibodies, the
vectors encoding the antibodies were expressed in
5 Chinese Hamster Ovary cells using the following
protocol.

A large scale plasmid DNA was purified using the
WIZARD® Maxipreps DNA Purification System (Promega
Catalog # A7421). The purified DNA was digested with
10 Ssp I and BspLU11 I, precipitated with ethanol once, and
resuspended in sterile TE.

Purified, endonuclease restricted plasmid DNA was
then introduced into Chinese hamster ovary (CHO)
dihydrofolate reductase minus DG44 cells using
15 electroporation. The electroporation technique used is
described below.

Approximately 1.6×10^8 CHO cells were spun in an
appropriate size sterile Corning tube for one minute at
1000 RPM. The media was removed and the cells were
20 washed in fifteen milliliters of sterile ice cold SBS
(sterile sucrose buffered solution is 272 mM sucrose, 7
mM sodium phosphate pH 7.4, 1 mM $MgCl_2$) and spun for 5
minutes at 1000 RPM. The SBS was removed and cells were
suspended using fresh ice cold sterile SBS at a cell
25 concentration of 1×10^7 cells were per ml and left on ice
for 15 minutes. The BTX 600 electroporator was turned
on and preset at 230 volts, with the maximum voltage
knobs being set at 500 volts/capacitance & resistance.
The capacitance was set at 400 microfaradays and the
30 resistance was set at 13 ohms (setting R1).

Plasmid DNA (4 μ g DNA or 2 μ g DNA) and 0.4 ml of
cells (4×10^6 cells) were then placed in BTX 0.4 ml
cuvettes (BTX Catalog # 620). The cells were shocked by
placing the cuvette into the BTX 600 stand and pressing

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the automatic charge & pulse button. Approximately 20 separate electroporations were performed with each mammalian expression plasmid.

After shocking, the cuvettes were left at ambient temperature for fifteen minutes. The cells and DNA were from each cuvette were resuspended in 20 ml of CHO-SSFMII containing no hypoxanthine or thymidine (Gibco BRL Catalog # 31033-012) to which HT supplement (100X supplement is 10 mM sodium hypoxanthine, 1.6 mM thymidine Gibco BRL Catalog # 11067-014) had been added. The cells from a single electroporated cuvette were then plated into a 96 well plates (200 μ l/well) and placed into a 37°C CO₂ incubator. Selection was started two or three days later by changing the media to the above media with the addition of 400 mg/ml of Geneticin® (G418, Gibco BRL Catalog # 10131-019). The cells were grown at 37°C and the cell media were changed every 3-5 days. After sixteen days G418 resistant clones appeared in the wells and the supernatant was assayed for antibody expression by ELISA. The highest expressing clones were then expanded individually. Monoclonal antibodies were purified as described below.

Immunoglobulin ELISA

Plates (Immulon 2, Dynatech Laboratories, Inc. Catalog # 011-010-3455) are coated overnight at 4°C with 200 ng unlabeled goat anti-human IgG antibody at 100 μ l/well. This is effected using twenty milliliters of unlabeled goat anti-human IgG/10 mls Coating Buffer/plate (Boehringer Mannheim Ab Catalog # 605 400). (1:500 dilution of ~ 1mg/ml stock.) The coating buffer is then removed from the plates and dried using a paper towel. One hundred microliters of a dilution buffer/well is then added.

Antibody solutions and standards (100 ng/ml - 2.5 ng/ml) are then added in duplicate at 100 μ l/well directly to the 100 ml dilution buffer. The antibody solutions and standards are contained in dilution
5 buffer. The resultant solutions are then incubated for at least 1 hour at 37°C.

After incubation, the contents of each plate are removed and the plates are washed with tap water five times. The plates are then dried on a paper towel.

10 After drying of the plates, a second antibody is then added at 100 μ l/well. This second antibody is either goat anti-human Kappa-HRPO: added at 1/10,000 dilution or 1 μ l Ab/10 mls dilution buffer/plate, available from Southern Biotechnology Associates, Inc.
15 Catalog # 2060-05 or a goat anti-human Lambda-HRPO; used at 1/20,000 dilution or 1 μ l Ab/20 mls dilution buffer/2 plates (available from Southern Biotechnology Associates, Inc. Catalog # 2070-05).

20 The antibody and contents of the plate are allowed to incubate for one hour at 37°C. After incubation the contents of each plate are removed. The plates are again washed five time with tap water, and the washed plates are dried. To the dried plates is then added HRPO substrate (TMB Microwell - two component) in an
25 amount of 100 μ l/well. (Five milliliters of TMB Peroxidase Substrate + five milliliters of Peroxidase Solution B/plate (Kirdgaard and Perry Labs, TMB Microwell two component reagents Catalog # 50-76-00).

30 The reaction is stopped by the addition of one hundred microliters of 2M H₂SO₄ to each well when the weakest standard (2.5 ng/ml) is visible over background. The optical density of wells in plates is then read using a plate reader, e.g., Molecular Devices Emax

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ELISA BUFFERS

5 Sodium Carbonate 0.8 gram / liter
Sodium Bicarbonate 1.55 gram / liter
Adjust pH to 9.5 with ~1ml 1N HCl

0.5% Nonfat Dry Milk in PBS plus 0.01% Thimerosal
10 (5 gm/L) (100 mg/L)

	<u>Standard</u>	<u>OD 450</u>	<u>OD450</u>	<u>Average</u>
15	100 ng/ml	0.805	0.876	0.841
	50 ng/ml	0.395	0.472	0.434
	25 ng/ml	0.213	0.252	0.233
	10 ng/ml	0.089	0.105	0.097
	5 ng/ml	0.054	0.055	0.055
	2.5 ng/ml	0.031	0.035	0.033
20	0 ng/ml	0.004	0.006	0.005

Appropriate dilution of stock AB (sterile filtered in normal saline, protein determination by OD) to give 1mg/ml

EXAMPLE:

Chimeric monkey/human anti-CD4 (CE9.1) is 4.18 mg/ml

24 μ l of above into 76 μ l Dilution Buffer is 1 mg/ml

50 μ l Stock Ab (1 mg/ml) into 450 μ l Dilution Buffer (D.B.) is 100 μ g/ml

50 μ l of above mixture into 450 μ l D.B. is 10 μ g/ml

200 μ l of above mixture into 1.8 mls D.B. is 1 μ g/ml

1 ml of above mixture into 9 mls D.B. is 100 ng/ml

*

5 ml of above mixture into 5 ml D.B. is 50 ng/ml *

5 ml of above mixture into 5 ml D.B. is 25 ng/ml *

4 ml of above mixture into 6 ml D.B. is 10 ng/ml *

5 ml of above mixture into 5 ml D.B. is 5 ng/ml *

5 ml of above mixture into 5 ml D.B. is 2.5 ng/ml *

* Standards used in the ELISA

Antibody purification by Protein A

Procedure

The culture supernatant is centrifuged to remove cells and debris. The centrifuge is then filtered through a 0.2 μ m filter. A protein A sepharose Fast flow column (recombinant protein A Sepharose Fast flow) (Pharmacia Biotech Catalog # 71-5000-09) is then prepared and equilibrated using PBS (pH 7.4).

The supernatant is loaded on the column at an appropriate flow rate (e.g., 2 ml/min). After loading, the column is washed with 10 column volume of PBS (pH 7.4). The antibody is eluted from the column using an elution buffer (0.2M acetic acid, 0.1 M glycine pH 3.5) at 1 ml/min flow rate. One milliliter fractions/tube

including 100 μ l of Tris are then collected. A spectrophotometer absorbance reading is then taken at 280 nm. The antibody fractions are then collected and dialyzed against PBS (pH 7.9) overnight. The dialysate
5 is then sterilized by filtration through a 0.22 μ m membrane and stored at -20°C.

Assay Results

The PRIMATIZED® human gamma-4 anti-human CD23 antibodies which are described *supra*, were purified and as-
10 sayed for induced IgE inhibitory activity *in vitro*. These results are contained in Figures 3 and 5. This was effected using the *in vitro* IL-4 IgE assay described *supra*.

These assay results surprisingly indicated that
15 both human gamma-4 anti-human CD23 antibodies were not as active as the corresponding primate anti-human CD23 antibodies, i.e., they did not significantly inhibit induced IgE production *in vitro*.

However, because primate 5E8 and p5E8G4P have a
20 potential asparagine linked glycosylation site in the heavy chain variable region, the effects of glycosylation at this site were investigated. (It was found that both these antibodies contain N-linked oligosaccharides at this site. (Data not shown.))
25 Therefore, in order to prevent glycosylation, the asparagine in the glycosylation site was changed to a lysine in order to eliminate carbohydrate addition. This mutated antibody was named p5E8G4PN-. Assay results demonstrated that this antibody behaved
30 identically to p5E8G4P in the IL-4 IgE assay (see Figure 3) and also exhibited an identical apparent affinity Kd for human CD23. (See Figure 4.) Therefore, these

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results indicated that the difference in IgE inhibition observed from the 5E8 gamma 4 PRIMATIZED® antibody in comparison to primate 5E8 antibody was not attributable to glycosylation differences.

5 The three primate antibodies (p5E8G4P, p5E8G4PN-, and pGG5G4P) were then expressed as human gamma-1 versions using substantially the same methodology. All three human gamma-1 anti-human CD23 antibodies, respectively designated p5E8G1, p5E8G1N- and pGG5G1,
10 were found to be active in the *in vitro* IL-4/IgE assay (Figures 3 and 5).

 p5E8G1 was found to be statistically more suppressive than p5E8G4P at a concentration of 0.3 µg/ml (P[T,t] one tail + 0.0055) and at 3 µg/ml (p[T<t] one
15 tail + 0.0019). In addition, p5E8G1N- is statistically more suppressive than p5E8G4PN- at both 0.3 µg/ml (p[T<t] one tail + 0.0392) and at 3 µg/ml (p[T<t] one tail + 0.0310) (Figure 3).

 Similarly, p6G5G1 completely inhibited induced IgE
20 production at 3 mg/ml, while p6G5G4P did not. (These results are in Figure 5).

 Thus, these results suggested that an active Fc region, in particular that of human gamma-1, is significant for induced IgE inhibition by anti-human CD23 anti-
25 bodies. These results also suggest that human anti-CD23 antibodies containing human gamma-3 constant regions will be significant for induced IgE inhibition because a gamma-3 constant region binds to the same FcγR receptors that gamma-1 binds to, and will likely mediate the same
30 result.

EXAMPLE 2

To confirm our hypothesis as to the involvement of the Fc effector portion in IgE inhibition of anti-human

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CD23 antibodies, a third primate antibody, designated 2C8, also shown to inhibit IgE in *in vitro*) was converted to a F(ab')₂. IgE inhibitory activity was determined using the same IL-4/IgE assay described previously.

Materials

The following materials were used in this example. ImmunoPure F(ab')₂ Preparation Kit (Pierce Catalog # 44888)

10 digestion buffer: 20 mM sodium acetate buffer, pH 4.5

0.1 M citric acid, pH 3.0 (adjust pH with NaOH)

0.1% sodium azide in water

dialysis tubing; 50,000 MW cutoff (Spectra Por

15 Catalog # 132 128)

shaking water bath capable of maintaining 37°C

polystyrene culture tubes, 17 x 100 mm (Fisher Catalog # 14-956-6B)

BCA protein assay (Pierce Catalog # 23224)

20 Centricon-50 concentrators (Amicon Catalog # 4225).

Equilibration of Immobilized Pepsin

0.25 milliliters of the 50% slurry of Immobilized Pepsin is added to a 17 x 100 mm test tube (0.125 ml of gel). Four milliliters of digestion buffer are then

25 added. The pepsin is then separated from the buffer using the serum separator. The buffer is then discarded and the wash procedure repeated using another four milliliters of buffer. The immobilized pepsin is then resuspended in 0.5 ml of digestion buffer.

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Preparation of Immobilized Protein A Column

Protein A AffinityPak® columns and ImmunoPure Binding and Elution buffers are brought to room temperature.

5 **Preparation of 2C8 F(ab')₂ Fragments**

2C8 F(ab')₂ fragments are prepared by methods well known in the antibody art. The inventors elected to use a commercially available kit, ImmunoPure F(ab')₂ Preparation Kit (Pierce Catalog # 44888), using the
10 manufacturer's protocols.

Ten milligrams of lyophilized 2C8 antibody were dissolved in one milliliter of a digestion buffer (20 mM sodium acetate buffer, pH 4.5). One milliliter of the antibody containing sample was then added to a tube con-
15 taining immobilized pepsin.

The antibody and immobilized pepsin were then incubated for four hours in a high speed shaker water bath at 37°C (at high speed), taking care to maintain the mixing constant during the incubation.

20 The resultant solubilized F(ab')₂ and Fc fragments and the undigested IgGs were then recovered from the immobilized pepsin gel using a serum separator. The crude digest is then decanted into a clean tube.

In order to enhance recovery of F(ab')₂ fragments,
25 the immobilized pepsin desirably is then washed with 1.5 of milliliters of the ImmunoPure IgG binding buffer. The wash is then added to the crude digest.

The antibody fragments were then recovered using a protein A column. This is effected by opening an
30 immobilized protein A column. Care is taken to avoid air bubbles from being drawn into the gel. The storage solution (which contains 0.02% sodium azide) is discarded.

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The immobilized protein A column was then equilibrated using twelve milliliters of binding buffer (contained in ImmunoPure Preparation Kit). The column was then transferred to a 17 x 100 mm test tube
5 contained in the kit (labeled "F(ab')₂") to collect eluate.

Three milliliters of the crude digest was then applied to a column and are allowed to flow completely into the gel. The use of AffinityPak™ columns is
10 desirable as these columns stop flowing automatically when the level reaches the top frit.

The column is then washed using six milliliters of binding buffer. The eluate which contains F(ab')₂ fragments was then collected. This eluate also contains
15 small Fc fragments that can no longer bind protein A (which are not bound to the Protein A column). However, the substantial portion thereof was eliminated by dialysis.

Dialysis was effected by taking the F(ab')₂ containing eluate and dialyzing the eluate against pH
20 7.4 phosphate buffered saline, using dialysis tubing with a molecular weight cut-off of 50,000 so as to eliminate the small Fc fragment containments (Spectra Pur. Catalog # 132 128).

This resulted in a F(ab')₂ fraction having an
25 optical density of 280 nm of 0.707 (6 ml). After dialysis and concentration with Centricon-50 concentrators (Amicon Catalog # 4225), the 2C8 F(ab')₂ product was assayed for protein content using a BCA
30 protein assay (Pierce Catalog # 23224). The protein content was found to be 3.76 mg per milliliter.

The 2C8 F(ab)₂'s were assayed for IgE inhibitory activity and were found to be substantially incapable of inhibiting IgE production in the same *in vitro* assays

described previously. These results are contained in Figure 6. In fact, the $F(ab')_2$ was found to antagonize the suppressive effects of induced IgE on the monoclonal antibody 2C8. These results are in Figure 7.

5

EXAMPLE 3

10 The PRIMATIZED® gamma 1 and gamma 4P versions of primate monoclonal 6G5 were both evaluated for their effect on inhibition of induced IgE production *in vivo* in the SCID mouse model described previously. p5E8G1N-
15 was found to be as efficient as primate 5E8 in inhibiting induced IgE. (See Figures 8 and 9). While neither primate 6G5 nor the primatized p6G5G4P were effective at inhibiting induced IgE *in vivo*, primatized p6G5G1 inhibited induced IgE production. (See Figures 9 and 10.) These results further substantiate our
20 conclusion that an active Fc region is significant to the ability of an anti-human CD23 antibody to effectively inhibit induced IgE production.

Proposed Mechanism

20 Although the results of the present invention conclusively demonstrate that an active Fc region is significant to the ability of an anti-human CD23 antibody to effectively inhibit induced IgE production, the molecular mechanism has not been elucidated.
25 However, several hypotheses may be made.

Firstly, it may be that the anti-CD23 antibodies of the present invention function to trigger a similar signal transduction pathway as is apparently triggered by cross-linking B cell receptor with anti-BCR gamma-1
30 antibodies. It has been hypothesized that the down-regulation of antigen receptor signaling in this case results from coligation of Fc gamma RII and surface Ig

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(B cell receptor) on the same B cell, leading to down regulation of Ig expression. (D'Ambrosia et al., 1995, *Science*, 268:293-297; Ono et al., 1997, *Cell* 90: 293-301). Indeed, CD23 is upregulated on the surface of IgE-expressing B cells, so it is possible that coligation of CD23 and the FcγRII-B receptor via the gamma-1 constant region results in a similar signal transduction pathway.

Alternatively, inhibition of induced IgE expression may be occurring through cross-linking of surface IgE on B cells at the same time as the Fc region interacts with an appropriate FcγR receptor on another cell type, i.e. a killer T cell, which may then "instruct" the B cell to undergo apoptosis, or lead to cell death in some other manner (i.e. phagocytosis). For instance, U.S. Patent No. 5,543,144, herein incorporated by reference, proposes a similar mechanism for inhibiting IgE-expressing B cells with anti-IgE antibodies. As discussed therein, antibodies of certain IgG subclasses, such as mouse IgG2a and human IgG1 and IgG3, can mediate ADCC (Antibody Dependent Cellular Cytotoxicity) carried out by certain Fc receptor-bearing phagocytic leukocytes. For example, OKT3, a mouse IgG2a monoclonal antibody specific for human T cell surface antigen (which was the first monoclonal antibody product approved by the FDA for marketing as a therapeutic agent) is used in patients to provide rapid depletion of T cells in the blood and to induce an immunosuppressed state (for kidney transplantation). (Russell, P. S. et.al, *Transpl. Proc.* 17:39-41 (1985)). OKT3, at a dosage of 5 mg/day/subject, can completely deplete circulating T cells. The monoclonal antibodies described in U.S. Patent 5,543,144, especially in the form of mouse gamma 2a antibodies or human or humanized

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antibodies bearing human gamma-1 or gamma-3 chains, are proposed to deplete IgE-expressing B cells by the ADCC mechanism in a similar manner.

But whatever the actual molecular mechanism for
5 inhibiting IgE expression using anti-CD23 antibodies may be, it is clear that the Fc region, and accordingly effector function, of anti-CD23 antibodies are important considerations. As discussed previously, since
10 complement is not present in the in vitro assays of the present invention, the phenomenon likely involves one or more FcγR receptors. Once the relevant molecular components are identified, as well as molecules internal to the cells involved which are involved in signal transduction pathways, it will be possible to design
15 other therapeutics which may also be used to inhibit induced IgE expression.

Utility

Because of their ability to effectively inhibit IgE production, the subject anti-human CD23 antibodies which
20 comprise human gamma-1 constant domains, and those that may also be constructed containing gamma-3, are effective in treating any disease wherein inhibition of IgE production is therapeutically desirable. Such diseases include by way of example allergic diseases,
25 autoimmune diseases and inflammatory disease.

Specific conditions which are potentially treatable by administration of the subject anti-CD23 human gamma-1/ gamma-3 constant domain containing antibodies include the following:

30 Allergic bronchopulmonary aspergillosis; Allergic rhinitis and conjunctivitis autoimmune hemolytic anemia; Acanthosis nigricans; Allergic contact dermatitis; Addison's disease; Atopic dermatitis; Alopecia areata; Alopecia universalis; Amyloidosis; Anaphylactoid

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purpura; Anaphylactoid reaction; Aplastic anemia; Angio-
edema, hereditary; Angioedema, idiopathic; Ankylosing
spondylitis; Arteritis, cranial; Arteritis, giant cell;
Arteritis, Takayasu's; Arteritis, temporal; Asthma;
5 Ataxia-telangiectasia; Autoimmune oophoritis; Autoimmune
orchitis; Autoimmune polyendocrine failure; Behcet's
disease; Berger's disease; Buerger's disease;
bronchitis; Bullous pemphigus; Candidiasis, chronic
mucocutaneous; Caplan's syndrome; Post-myocardial
10 infarction syndrome; Post-pericardiotomy syndrome;
Carditis; Celiac sprue; Chagas's disease;
Chediak-Higashi syndrome; Churg-Strauss disease; Cogan's
syndrome; Cold agglutinin disease; CREST syndrome;
Crohn's disease; Cryoglobulinemia; Cryptogenic fibrosing
15 alveolitis; Dermatitis herpetiformis; Dermatomyositis;
Diabetes mellitus; Diamond-Blackfan syndrome; DiGeorge
syndrome; Discoid lupus erythematosus; Eosinophilic
fasciitis; Episcleritis; Drythema elevatum diutinum;
Erythema marginatum; Erythema multiforme; Erythema
20 nodosum; Familial Mediterranean fever; Felty's syndrome;
Fibrosis pulmonary; Glomerulonephritis, anaphylactoid;
Glomerulonephritis, autoimmune; Glomerulonephritis,
post-streptococcal; Glomerulonephritis,
post-transplantation; Glomerulopathy, membranous;
25 Goodpasture's syndrome; Graft-vs.-host disease;
Granulocytopenia, immune-mediated; Granuloma annulare;
Granulomatosis, allergic; Granulomatous myositis;
Grave's disease; Hashimoto's thyroiditis; Hemolytic
disease of the newborn; Hemochromatosis, idiopathic;
30 Henoch-Schoenlein purpura; Hepatitis, chronic active and
chronic progressive; Histiocytosis_X; Hypereosinophilic
syndrome; Idiopathic thrombocytopenic purpura; Job's
syndrome; Juvenile dermatomyositis; Juvenile rheumatoid
arthritis (Juvenile chronic arthritis); Kawasaki's

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disease; Keratitis; Keratoconjunctivitis sicca;
Landry-Guillain-Barre-Strohl syndrome; Leprosy,
lepromatous; Loeffler's syndrome; lupus; Lyell's
syndrome; Lyme disease; Lymphomatoid granulomatosis;
5 Mastocytosis, systemic; Mixed connective tissue disease;
Mononeuritis multiplex; Muckle-Wells syndrome;
Mucocutaneous lymph node syndrome; Mucocutaneous lymph
node syndrome; Multicentric reticulohistiocytosis;
Multiple sclerosis; Myasthenia gravis; Mycosis
10 fungoides; Necrotizing vasculitis, systemic; Nephrotic
syndrome; Overlap syndrome; Panniculitis; Paroxysmal
cold hemoglobinuria; Paroxysmal nocturnal
hemoglobinuria; Pemphigoid; Pemphigus; Pemphigus
erythematosus; Pemphigus foliaceus; Pemphigus vulgaris;
15 Pigeon breeder's disease; Pneumonitis, hypersensitivity;
Polyarteritis nodosa; Polymyalgia rheumatic;
Polymyositis; Polyneuritis, idiopathic; Portuguese
familial polyneuropathies; Pre-eclampsia/eclampsia;
Primary biliary cirrhosis; Progressive systemic
20 sclerosis (Scleroderma); Psoriasis; Psoriatic arthritis;
Pulmonary alveolar proteinosis; Pulmonary fibrosis,
Raynaud's phenomenon/syndrome; Reidel's thyroiditis;
Reiter's syndrome, Relapsing polychondritis; Rheumatic
fever; Rheumatoid arthritis; Sarcoidosis; Scleritis;
25 Sclerosing cholangitis; Serum sickness; Sezary syndrome;
Sjogren's syndrome; Stevens-Johnson syndrome; Still's
disease; Subacute sclerosing panencephalitis;
Sympathetic ophthalmia; Systemic lupus erythematosus;
Transplant rejection; Ulcerative colitis;
30 Undifferentiated connective tissue disease; Urticaria,
chronic; Urticaria, cold; Uveitis; Vitiligo;
Weber-Christian disease; Wegener's granulomatosis;
Wiskott-Aldrich syndrome.

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Of these, the preferred indications treatable or presentable by administration of anti-CD23 antibodies include allergic rhinitis and conjunctivitis, atopic dermatitis; eczema; Job's syndrome, asthma; allergic conditions; and chronic inflammatory diseases and conditions, including CLL chronic lymphocytic leukemia, typically characterized by high levels of membrane CD23 and high circulating levels of sCD23 (Sarfart et al., Blood 71: 94-98 (1988)).

The amount of antibody useful to produce a therapeutic effect can be determined by standard techniques well known to those of ordinary skill in the art. The antibodies will generally be provided by standard technique within a pharmaceutically acceptable buffer, and may be administered by any desired route. Because of the efficacy of the presently claimed antibodies and their tolerance by humans it is possible to administer these antibodies repetitively in order to combat various diseases or disease states within a human.

One skilled in the art would be able, by routine experimentation, to determine what an effective, non-toxic amount of antibody would be for the purpose of effecting allergic diseases and inflammatory conditions. Generally, however, an effective dosage will be in the range of about 0.05 to 100 milligrams per kilogram body weight per day.

The antibodies of the invention may be administered to a human or other animal in accordance with the aforementioned methods of treatment in an amount sufficient to produce such effect to a therapeutic or prophylactic degree. Such antibodies of the invention can be administered to such human or other animal in a conventional dosage form prepared by combining the

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antibody of the invention with a conventional pharmaceutically acceptable carrier or diluent according to known techniques. It will be recognized by one of skill in the art that the form and character of the pharmaceutically acceptable carrier or diluent is dictated by the amount of active ingredient with which it is to be combined, the route of administration and other well-known variables.

The route of administration of the antibody of the invention may be oral, parenteral, by inhalation or topical. The term parenteral as used herein includes intravenous, intramuscular, subcutaneous, rectal, vaginal or intraperitoneal administration. The intravenous form of parenteral administration is generally preferred.

The daily parenteral and oral dosage regimens for employing compounds of the invention to prophylactically or therapeutically induce immunosuppression will generally be in the range of about 0.05 to 100, but preferably about 0.5 to 10, milligrams per kilogram body weight per day.

The antibody of the invention may also be administered by inhalation. By "inhalation" is meant intranasal and oral inhalation administration. Appropriate dosage forms for such administration, such as an aerosol formulation or a metered dose inhaler, may be prepared by conventional techniques. The preferred dosage amount of a compound of the invention to be employed is generally within the range of about 10 to 100 milligrams.

The antibody of the invention may also be administered topically. By topical administration is meant non-systemic administration and includes the application of an antibody (or fragment thereof) compound of the

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invention externally to the epidermis, to the buccal cavity and instillation of such an antibody into the ear, eye and nose, and where it does not significantly enter the blood stream. By systemic administration is meant oral, intravenous, intraperitoneal and intramuscular administration. The amount of an antibody required for therapeutic or prophylactic effect will, of course, vary with the antibody chosen, the nature and severity of the condition being treated and the animal undergoing treatment, and is ultimately at the discretion of the physician. A suitable topical dose of an antibody of the invention will generally be within the range of about 1 to 100 milligrams per kilogram body weight daily.

Formulations

While it is possible for an antibody or fragment thereof to be administered alone, it is preferable to present it as a pharmaceutical formulation. The active ingredient may comprise, for topical administration, from 0.001% to 10% w/w, e.g., from 1% to 2% by weight of the formulation, although it may comprise as much as 10% w/w but preferably not in excess of 5% w/w and more preferably from 0.1% to 1% w/w of the formulation.

The topical formulations of the present invention, comprise an active ingredient together with one or more acceptable carrier(s) therefor and optionally any other therapeutic ingredients(s). The carrier(s) must be "acceptable" in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof.

Formulations suitable for topical administration include liquid or semi-liquid preparations suitable for penetration through the skin to the site of where treat-

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ment is required, such as liniments, lotions, creams, ointments or pastes, and drops suitable for administration to the eye, ear or nose.

5 Drops according to the present invention may comprise sterile aqueous or oily solutions or suspensions and may be prepared by dissolving the active ingredient in a suitable aqueous solution of a bactericidal and/or fungicidal agent and/or any other suitable preservative, and preferably including a
10 surface active agent. The resulting solution may then be clarified by filtration, transferred to a suitable container which is then sealed and sterilized by autoclaving or maintaining at 90°-100°C for half an hour. Alternatively, the solution may be sterilized by
15 filtration and transferred to the container by an aseptic technique. Examples of bactericidal and fungicidal agents suitable for inclusion in the drops are phenylmercuric nitrate or acetate (0.002%), benzalkonium chloride (0.01%) and chlorhexidine acetate
20 (0.01%). Suitable solvents for the preparation of an oily solution include glycerol, diluted alcohol and propylene glycol.

25 Lotions according to the present invention include those suitable for application to the skin or eye. An eye lotion may comprise a sterile aqueous solution optionally containing a bactericide and may be prepared by methods similar to those for the preparation of drops. Lotions or liniments for application to the skin may also include an agent to hasten drying and to cool
30 the skin, such as an alcohol or acetone, and/or a moisturizer such as glycerol or an oil such as castor oil or arachis oil.

Creams, ointments or pastes according to the present invention are semi-solid formulations of the

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active ingredient for external application. They may be made by mixing the active ingredient in finely-divided or powdered form, alone or in solution or suspension in an aqueous or non-aqueous fluid, with the aid of

5 suitable machinery, with a greasy or non-greasy basis. The basis may comprise hydrocarbons such as hard, soft or liquid paraffin, glycerol, beeswax, a metallic soap; a mucilage; an oil of natural origin such as almond, corn, arachis, castor or olive oil; wool fat or its
10 derivatives, or a fatty acid such as stearic or oleic acid together with an alcohol such as propylene glycol or macrogels. The formulation may incorporate any suitable surface active agent such as an anionic, cationic or non-ionic surface active such as sorbitan
15 esters or polyoxyethylene derivatives thereof. Suspending agents such as natural gums, cellulose derivatives or inorganic materials such as siliceous silicas, and other ingredients such as lanolin, may also be included.

20 It will be recognized by one of skill in the art that the optimal quantity and spacing of individual dosages of an antibody or fragment thereof of the invention will be determined by the nature and extent of the condition being treated, the form, route and site of
25 administration, and the particular animal being treated, and that such optimums can be determined by conventional techniques. It will also be appreciated by one of skill in the art that the optimal course of treatment, i.e., the number of doses of an antibody or fragment thereof
30 of the invention given per day for a defined number of days, can be ascertained by those skilled in the art using conventional course of treatment determination tests.

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Without further elaboration, it is believed that one skilled in the art can, using the preceding description, utilize the present invention to its fullest extent. The following are, therefore, to be construed as merely illustrative examples and not a limitation of the scope of the present invention in any way.

Capsule Composition

A pharmaceutical composition of this invention in the form of a capsule is prepared by filling a standard two- piece hard gelatin capsule with 50 mg. of an antibody or fragment thereof of the invention, in powdered form, 100 mg. of lactose, 32 mg. of talc and 8 mg. of magnesium stearate.

Injectable Parenteral Composition

A pharmaceutical composition of this invention in a form suitable for administration by injection is prepared by stirring 1.5k by weight of an antibody or fragment thereof of the invention in 10k by volume propylene glycol and water. The solution is sterilized by filtration.

Ointment Composition

Antibody or fragment thereof of the invention 10 g.
White soft paraffin to 100.0 g.
The antibody or fragment thereof of the invention is dispersed in a small volume of the vehicle to produce a smooth, homogeneous product. Collapsible metal tubes are then filled with the dispersion.

Topical Cream Composition

Antibody or fragment thereof of the invention 1.0g.

Polawax GP 200 20.0 g.
Lanolin Anhydrous 2.0 g.
White Beeswax 2.5 g.
Methyl hydroxybenzoate 0.1 g.
5 Distilled Water to 100.0 g.

The polawax, beeswax and lanolin are heated together at 60°C. A solution of methyl hydroxybenzoate is added and homogenization is achieved using high speed stirring. The temperature is then allowed to fall to 10 SOOC. The antibody or fragment thereof of the invention is then added and dispersed throughout, and the composition is allowed to cool with slow speed stirring.

Topical Lotion Composition

Antibody or fragment thereof of the invention 1.0g.
15 Sorbitan Monolaurate 0.6 g. Polysorbate 20 0.6 g.
Cetostearyl Alcohol 1.2 g. Glycerin 6.0 g.
Methyl Hydroxybenzoate 0.2 g.

Purified Water B.P. to 100.00 ml. (B.P. = British Pharmacopeia)

20 The methyl hydroxybenzoate and glycerin are dissolved in 70 ml. of the water at 75°C. The sorbitan monolaurate, polysorbate 20 and cetostearyl alcohol are melted together at 75°C and added to the aqueous solution. The resulting emulsion is homogenized,
25 allowed to cool with continuous stirring and the antibody or fragment thereof of the invention is added as a suspension in the remaining water. The whole suspension is stirred until homogenized.

Eye Drop Composition

30 Antibody or fragment thereof of the invention 0.5g.
Methyl Hydroxybenzoate 0.01 g.
Propyl Hydroxybenzoate 0.04 g.

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Purified Water B.P. to 100.00 ml.

The methyl and propyl hydroxybenzoates are dissolved in 70 ml. purified water at 75°C and the resulting solution is allowed to cool. The antibody or fragment thereof of the invention is then added, and the solution is sterilized by filtration through a membrane filter (0.022 Am pore size), and packed aseptically into suitable sterile containers.

Composition for Administration by Inhalation

10 For an aerosol container with a capacity of 15-20 ml: mix 10 mg. of an antibody or fragment thereof of the invention with 0.2-0.5k of a lubricating agent, such as polysorbate 85 or oleic acid, and disperse such mixture in a propellant, such as freon, preferably in a
15 combination of (1,2 dichlorotetrafluoroethane) and difluorochloromethane and put into an appropriate aerosol container adapted for either intranasal or oral inhalation administration. Composition for
20 Administration by Inhalation For an aerosol container with a capacity of 15-20 ml: dissolve 10 mg. of an antibody or fragment thereof of the invention in ethanol (6-8 ml.), add 0.1-0.2k of a lubricating agent, such as polysorbate 85 or oleic acid; and disperse such in a propellant, such as freon, preferably in combination of
25 (1-2 dichlorotetrafluoroethane) and difluorochloromethane, and put into an appropriate aerosol container adapted for either intranasal or oral inhalation administration.

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Parenteral Administrable Antibody Compositions

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The antibodies and pharmaceutical compositions of the invention are particularly useful for parenteral administration, i.e., subcutaneously, intramuscularly or intravenously. The compositions for parenteral administration will commonly comprise a solution of an antibody of the invention or a cocktail thereof dissolved in an acceptable carrier, preferably an aqueous carrier. A variety of aqueous carriers may be employed, e.g., water, buffered water, 0.4k saline (normal saline), 0.3% glycine, and the like. The use of normal saline is preferred. These solutions are sterile and generally free of particulate matter. These solutions may be sterilized by conventional, well-known sterilization techniques. The compositions may contain pharmaceutically acceptable auxiliary substances as required to approximate physiological conditions such as pH adjusting and buffering agents, etc. The concentration of the antibody or fragment thereof of the invention in such pharmaceutical formulation can vary widely. Such concentrations will be selected primarily based on fluid volumes, viscosities, etc., according to the particular mode of administration selected. Generally suitable intravenous concentrations range from about one to one hundred milligrams per milliliter.

Thus, a pharmaceutical composition of the invention for intravenous injection could comprise 10 mL normal saline containing 40-50 mg of an anti-human CD23 antibody of the invention. Methods for preparing parenterally administrable compositions are well-known or will be apparent to those skilled in the art, and are described in more detail in, for example, *Remington's Pharmaceutical Science*, 15th ed., Mack Publishing

Company, Easton, Pennsylvania, hereby incorporated by reference herein.

5 The antibodies of the invention can be lyophilized for storage and reconstituted in a suitable carrier prior to use. This technique has been shown to be effective with conventional immune globulins and art-known lyophilization and reconstitution techniques can be employed.

10 Depending on the intended result, the pharmaceutical composition of the invention can be administered for prophylactic and/or therapeutic treatments. In therapeutic application, compositions are administered to a patient already suffering from a disease, in an amount sufficient to cure or at least
15 partially arrest the disease and its complications. In prophylactic applications, compositions containing the present antibodies or a cocktail thereof are administered to a patient not already in a disease state to enhance the patient's resistance.

20 Single or multiple administrations of the pharmaceutical compositions can be carried out with dose levels and pattern being selected by the treating physician. In any event, the pharmaceutical composition of the invention should provide a quantity of the
25 subject anti-CD23 antibodies sufficient to effectively treat the patient.

It should also be noted that the antibodies of this invention may be used for the design and synthesis of either peptide or non-peptide compounds (mimetics) which
30 would be useful in the same therapy as the antibody. See, e.g., Saragovi et al., *Science*, 253:792-795 (1991).

From the foregoing, it will be appreciated that, although specific embodiments of the invention have been described herein for purposes of illustration, various

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modifications may be made without diverting from the scope of the invention. Accordingly, the invention is not limited by the appended claims.

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